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online in File 415

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(Files 799, 772, 771)

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))) of new databases, price changes, etc.

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#### File 1:ERIC 66-93/JAN

Set Items Description

?5 155,50,55,144,76

C4feb93 11:16:05 User219509 Session D70.1

00.29 0.038 Mrs File1

60.29 Estimated most File!

\$0.09 DIRLNET

\$0.38 Estimated cost this search

t0.28 Esticated total session cost 0.009 Hrs.

SYSTEM: OS - DIALES CheSearch

File 155: MEDLINE 1986-1993/MAR (9303W4)

File 50:CAB Abstracts 1984-1993/Jan

(c) 1993 CAB International

\*\*FILEC50: COPR. 1990 CABI. See HELPNEWS50 for CAB training schedule.

File 55:BIOSIS PREVIEWS 85-93/JAN 8A9504:BARRM4404

(C. BIOSIS 1993)

\*\*FILE 55: Biosystepatic Codes (BC=) for viruses have changed for 1993.

Type ?NEWSSS for more information and a complete list of the new modes.

File 144:PASCAL 1973 - 1992 DEC

(C. INIST/CNRE 1992)

\*\*FILE144: Ligit probleg: see ?news144

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File 76:LIFE SCIENCES COLLECTION 70-92/DEC

(Copr. Capbridge Scientific Abs.)

Set Items Description

?b 155, 50, 55, 144, 72, 350, 351

04feb93 11:16:36 User219529 Session D70.2

60.03 0.031 Hrs File155

\$0.03 Estimated cost File155

00.06 0.001 Hrs File50

\$0.C5 Estimated cost File50

\$0.10 Estimated cost File55

00.26 0.001 Hrs File144

\$0.05 Estimated cost Fila144

\$0.09 0.001 Krs File76

60.09 Estimated cost File75 OneSearch, 5 files, C.028 Mrs FileCS

\$0.09 DIGLNET

\$0.43 Estimated cost this search

\$0.61 Estimated total session cost 0.016 Was.

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SYSTEM:US - DIALOS CheSearch
File 155:MEDLINE 1964-1992/MAR (9303W4)
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File 50:CAR Abstracts 1984-1993/Jan (c) 1993 CAB International

emFILE050: CEPR. 1990 CABI. See MELPNEWSES for CAB training schedule.

File 55:BIOSIS PREVIEWS 85-93/JAN BA9504:BARRM4404

(C. BIOSIS 1993)

##FILE 55: Biosystematic Codes (BC=) for viruses have changed for 1993.

Type ?NEWSSS for more inforcation and a complete list of the new codes.

File 144: PASCAL 1973 - 1992 DEC

(C. INIST/CNRS 1992)

\*\*FILE144: Limit problem: see ?news144

Update 9301 has been temporarily backed off due to duplicate records

File 72:EMBASE (EXCERPTA MEDICA) 85-93/ISBS5

(CCPR. ESP BV/EM 1993)

##FILE 72: Truncate ENTREE Codes (e.g. DC=C1.120?) for cooplete ##retrieval. The 9245 update has been added.

File 350:Derwent World Patents Index

1053-1980, EQUIVALENTS THRU DW=9247

@GFILE35C: Format 9 includes the expanded patent table.
Preformatted

REPORTs are available. Type ?FMT3E0, ?NEMS350, ?RATES350 for pore info.

File 351:DERWENT WORLD PATENTS INDEX-LATEST

1981+;DW=9251,UA=9239,UM=9208

\*\*FILE351: Format 9 includes the expanded patent table.
Preferented

REPORTs are available. Type ?FMT351, ?NEWS351, ?RATES351 for

core info.

Set Items Description

?s antivenin

SI 591 ANTIVENIN

?s s1 and (f(ab) or fab fragment or f(ab)2)

591 S1

C F(AB)

4 FAS FRASMENT

0 F(AB)2

SS 0 SI AND (F(AB) OR FAB FRAGMENT OR F(AB)2)

?s si and ( f(ab) or fab(w)fragment or fab or fab2 )

591 51

@ F(AB)

19255 FAB

133728 FRASKENT

2039 FAR (H) FRASMENT

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78 FAR2
     53
              2 SI AND ( F(AB) OR FAB(W) FRAGMENT OR FAB CR FAB2
?t s3/3/1,2
 3/3/1
          (Item 1 from file: 155)
06089508 87063508
 Immunotherapy in the poisoned patient. Overview of present
applications
and future trends.
 Sullivan JB Jr
 Med Toxicol Jan-Feb 1986, 1 (1) p47-50, ISSN 0112-5966
Journal Code: MDT
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE; REVIEW
 3/3/2
          (Item 1 from file: 144)
 10276259 PASCAL No.: 92-0482170
 An affinity purified ovine antivence for the treatment of
Vipera berus
envenoming
 SMITH D C; REDDI K R; LAING G; THEAKSTON R G D; LANDON J
  Therapeutic Antibodies Inc., St. Bartholomew's hosp. cadical
school,
London, United Kingdom
  Journal: Toxicon: (Oxford), 1992, 30 (8) 865-871
 Language: English
?ds
Set
       Items Description
S1
         591 ANTIVENIN
           0 SI AND (F(AB) OR FAB FRAGMENT OR F(AB)2)
           2 SI AND (F(AB) OR FAB(W)FRAGMENT OR FAB OR FAB2 )
?s venca
     54 23262 VENOM
?s fab or fab2
          19255 FAB
             78 FAB2
          19302 FAB OR FAB2
?s s4 and s5
          23862 $4
             72 S4 AND S5
282
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)))Duplicate detection is not supported for File 35%.)))Duplicate detection is not supported for File 351.

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    Records from unsupported files will be retained in the RD set.

...examined 50 records (50) ...completed examining records
     S7
            40 RD (unique items)
?t 57/3/1-40
 7/3/1
           (Item 1 from file: 155)
          93016233
68306233
  Toxicity associated with the formation and clearance of immune
complexes
           antitumour monoclonal antibodies and syngeneic
between
anti-idiotypic
antibodies in mice.
 Pimm MV; Gribben SJ
  Cancer Research Campaign Laboratories,
                                                  University
Nottingham, UK.
  J Cancer Res Clin Carol (SERMANY) 1992, 119 (1) p41-5, ISSN
0171-5216
Journal Code: HLS
 Languages: ENGLISH
  Document type: JCURNAL ARTICLE
           (Item 2 from file: 155)
 7/3/2
28259347 92397347
  An affinity purified ovine antivenous for the treatment of
Vipera berus
envenoming.
  Smith DC; Reddi KR; Laing G; Theakston RG; Landon J
  Therapeutic Antibodies
                            Inc., St. Bartholomew's Hospital
Medical School,
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4

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London, U.K.
                     Aug 1992, 30 (8) p865-71, ISSN 0041-0101
 Toxicon (ENGLAND)
Journal Code: WWT
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/3
           (Item 3 from file: 155)
08248192 92386192
 Limited proteclysis of human and rabbit immunoglobulins by
snake vencss
produces Fab-like fragments.
 Assakuya MT; Mandelbaum FR
 Servico de Bioquisica, Instituto Butantan, Sao Paulo, Brasil.
 Eraz J Med Biol Res (BRAZIL) 1990, 23 (12) p1233-5, ISSN
0100-879X
Journal Code: BOF
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/4
           (Item 4 from file: 155)
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07995510 92033510

.. . .

anticoagulant after allogeneic bone marrow transplantation. Morio S; Ch H; Hirasawa A; Astsuka N; Nakasura H; Asai T; Yoshida S; Ito Department of Internal Second Medicine, School of Medicine, Chiba University, Japan. 1991, 3 (2) Transplant (ENGLAND) Aug Bone Marro*a* p147-9, ISSN 0268-2369 | Journal Code: BCN Languages: ENGLISE Document type: JOURNAL ARTICLE 7/3/5 (Item 5 from file: 155) 07885296 92023296 enzymes of crotalid Neutralization of kinin-releasing venozs by monospecific and polyspecific antivenoms. Bailey GS; al-Joufi A; Rawat S; Smith DC Department of Chemistry and Biological Chemistry, University of Essex, Colchester, U.K. 1991, 29 (6) p777-81, ISSN 0041-0101 Journal Toxicon Code: VWT Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/6 (Item 6 from file: 155) 07804536 91323536 Crystallization and preliminary X-ray diffraction data of the Fab fragment of a conoclonal antibody against apazin, a bee venom meurotoxia. Devaux C; Defendini ML; Alzar PM; Abergel C; Granier C; Fontecilla-Camps Laboratoire de Biochimie, CNRS URA 1455, Faculte de Medecine-Nord, Marseille, France. FEBS Lett Jul 29 1991, 286 (1-2) p64-6, ISSN 0014-5793 Journal Code: EUH Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/7 (Item 7 from file: 155) **07552443** 91071443 A new small myotoxin from the venom of the prairie mattlesnake (Crotalus viridis viridis). Griffin PR; Rind SD California Institute of Technology, Division of Biology,

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Jaurnal Cade: EUM
 Languages: EMGLISH
 Document type: JOURNAL ARTICLE
7/3/8
          (Item 8 from file: 155)
07523655 91042655
 Complement killing of human neuroblastoma cells: a cytotoxic
antibody and its F(ab*)2-cobra venom factor conjugate
are equally
cytotoxic.
 Juhl H; Petrella EC; Cheung NK; Bredehorst R; Vogel CW
 Department of Biochemistry and Molecular Biology, Georgetown
University,
Washington, DC 20007.
 Mol Immunol Det 1990, 27 (10) p957-64, ISSN 0161-5890
Journal Code: NG1
 Contract/Grant No.: CA 35525; CA 01039
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
          (Item 9 from file: 155)
7/3/9
27476502 90385500
 Cleavage of immunoglobulins by mosjeni protease A, from
the venom of
Bothrops moojeni.
 Assakura MT; Mandelbaum FR
 Servico de Bioquisica, Instituto Butantan, Sao Paulo, Brasil.
 Toxicco 1990, 28 (6) p734-6, ISSN 0041
-0101 Journal Code: VWT
 Languages: ENGLISH
  Document type: JOURNAL ARTICLE
 7/3/10
           (Item 10 from file: 155)
07060610 89362510
  Immunotoxicotherapy: present status and future trends.
 Scherrmann JM; Terrien N; Urtizberea M; Pierson P; Denis H;
Bourre JM
  Insera U.25, Hopital Fernand Widal, Paris, France.
  J Toxical Clin Toxical 1989, 27 (1-2) p1-35, ISSN 0731-3910
Journal Code: KAN
  Languages: ENSLISH
  Document type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
 7/3/11
           (Item 11 from file: 155)
05939841 39291841
 Molecular recognition in the activation of human blood
coagulation factor
  Chattopadhyay A; Fair DS
  Department of Bischemistry, University of Texas Health
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75710.
 Journal Codes HIV
 Contract/Grant No.: HL 39040
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
          (Item 12 from file: 155)
7/3/12
28935725 89267705
 Purification of F(ab')2 anti-snake venom by caprylic acid: a
fast method
for obtaining IgS fraggents with high neutralization activity,
purity and
yield.
 dos Santos MC; D'Imperio Lima MR; Furtado GC; Colletto GM;
Kipnis TL;
Dias da Silva W
 Fundação Ezequiel Dias, Universidade Federal de Minas Gerais,
Brazil.
           1989, 27 (3) p297-303,
                                    ISSN 0041-0101
                                                    Journal
  Toxicon
Code: VWT
 Languages: ENGLISH
 Document type: JCURNAL ARTICLE
7/3/13
           (Item 13 from file: 155)
06869252 89171252
 Neutralization of lethal potency and inhibition of enzymatic
a phospholipase A2 neurotoxin, crotoxin, by non-precipitating
antibodies
(Fab).
 Chouret 7; Jiang MS; Radvanyi F; Cwmby C; Bon C
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Laboratoire des Vezins, Institut Pasteur, Paris, France.
 FEBS Lett Feb 13 1989, 244 (1) p167-73, ISSN 0014-5793
Journal Code: EUH
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/14
            (Item 14 from file: 155)
06525042 86170042
 Decay-accelerating
                        factor
                                   protects
                                                husan
                                                           tumor
cells from
complement-mediated cytotoxicity in vitro.
  Chaung NK; Walter EI; Saith-Mensah WH; Ratnoff WD; Tykocinski
ML: Medof
 Department of Pediatrics, Mesorial Sloan Kettoring Cancer
Centar, New
York 12321.
                 Apr 1988, 61 (4) p1122-9, ISSN 0021-9730
  J Clin Invest
Journal Code: MS7
  Contract/Grant No.: C9-39320, AI-24220; AI-22030
  Languages: ENGLISH
  דר בארל לייבה, זמיימיותו המדדמים
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7/3/15
           (Item 15 fyom file: 155)
06353393 88203393 Block of synaptic transmission in insect CNS by toxins from
the venor of
the wasp Megascolia flavifrons (Fab.).
 Piek T; Hue B; Mony L; Nakajima T; Galhate M; Yasuhama T
  Department of Pharmacology, University of Amsterdam, The
Netherlands.
 Comp Biochem Physiol [C3 1987, 87 (2) p287-95,
                                                            ISSN
6742-8413
Journal Code: DNX
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/16
           (Item 16 from file: 155)
05089508 87063508
  Immunotherapy in the poisoned patient. Overview of present
applications
and future trends.
 Sullivan JB Jr
 Med Toxicol Jan-Feb 1985, 1 (1) p47-50, ISSN 0112-5955
Journal Code: MD?
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE; REVIEW
           (Item 17 from file: 155)
7/3/17
06035955 87009955
  Defining the role of complement in experimental pemphigus
vulgaris in
sice.
  Anhalt SJ; Till SO; Diaz LA; Labib RS; Fatel HP; Eaglstein NF
  J Impunol Nev 1 1986, 137 (9) p2835-40, ISSN 0022-1757
Journal Code: IFB
 Contract/Grant No.: R23-AM32490; R23-AM32279; R21-AM32599; +
  Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/19
           (Item 18 from file: 155)
CE549020 85265020
 Role of complement and the Fc portion of immunoglobulin S in
iccunity to
Venezuelan
                              encephalomyelitis
                                                      virus
                 equine
           with
infection
glycoprotein-specific monoclonal antibodies.
 Mathews JH; Roehrig JT; Trent DM
 J Virol Sep 1985, 55 (3) p594-600, ISSN 0022-538X Journal
Code: KCV
  Languages: ENGLISH
  Document type: JOURNAL ARTICLE
7/3/19
           (Item 19 from file: 155)
25534749 85150749
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SHEET STREET SHEETEN CONTRACT STREET hasaatological changes in the rabbit by pethylpredmissions, F(ab1)2 fraggents and rosparinio acid. Bult H; Herman AS; Rampart M Pr J Pharmacel Feb 1985, 84 (2) p317-27, ISt s7/3/9-42 SN 2027-1188 Journal Code: BCC Languages: ENGLICH Document type: JOURNAL ANTICLE 7/3/22 (Itam 20 from file: 155) 05412032 85026032 Magascoliakinin, a bradykinin-like compound in the venom of Megasoclia flavifrons Fab. (Hymenoptera: Scoliidae). Piek T; Mantel P; Van Ginkel CJ Comp Biochem Physicl [C] 1984, 78 (2) p473-4, **@742-9413** Journal Code: DNX Languages: ENSLIGH Document type: JCURNAL ARTICLE 7/3/21 (Item 21 from file: 155) 04706845 82249845 icsunoglobulins accelerate Naurotoxin-specific dissociation of the neurotoxin-acetylcholine receptor complex. Boulain JC; Menez A Science Aug 20 1982, 217 (4561) p732-3, ISEN 0036-8075 Journal Code: UJ7 Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/22 (Item 22 from file: 155) 04228391 81056391 Inhibition of biological activity of souse beta-nerve growth factor by sonoclonal antibody. Warren SL; Fanger M; Neet KE Science Nov 21 1980, 210 (4472) p912-2, ISSN 2036-6075 Journal Code: UJ7 Contract/Grant No.: A110148; CR27915 Languages: EMCLISH Document type: JOURNAL ARTICLE 7/3/23 (Itan 23 from file: 155) 04183515 81011516 Issunopharkacological approach to Foressan shock.

Nagai M; Kurizoto Y; Koda A

Document type: JOURNAL ARTICLE

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Languages: ENGLIGH

7/2/01

Microbiol Issunol 1980, 24 (7) p649-55, Journal Osdo: MX7

24263395 88174695 Impune retention: immunological requirements for Taintaining an easily dagradeble antigen in vivo. Yew JG; Mandel TE; Miller GA Aust J Exp Biol Med Sci Rug 1979, 57 (4) p401-14, ISEN 0004-945X Journal Code: 9FW Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/25 (Item 25 from file: 155) 23671494 79248494 Immunological studies on pancreatic phospholipase A2. Antigenic characterization of the NH2-targinal region. Meijer M; Meddens MJ; Dijksan R; Slotboom AJ; de Haas EH J Biol Chem Dec 10 1978, 253 (23) p2554-9, ISSN 0321-9258 Journal Code: HIV Languages: ENGLISH Document type: JCURNAL ARTICLE (Item 26 from file: 155) 7/3/26 02909946 76090946 nephrotoxic nephritis. I. The role of Guinea-pig complement and polymorphonuclear leucocytes and the effect of antibody subclass and fragments in the heterologous phase. Simpson IJ; Amos N; Evans DJ; Thomson NM; Peters DX Clin Exp Immunel Mar 1975, 19 (3) p499-511, ISSN 2009-9104 Journal Code: DD7 Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/27 (Ites 1 from file: 55) 9149815 BIOSIS Number: 93134315 MOVEL QUATERNARY AXMONIUM SALT-CONTAINING POLYAMINES FROM THE AGELENOPSIS-AFERTA FUNNEL-WEB SPIDER JASYS V J; KELBAUGH P R; MASON D M; PHILLIPS D; ROSMACK K J; FORMAN J I; SACCOMAND N A; STRCH J S; VOLKMANN R A CENTRAL RESEARCH DIVISION, PFIZER INC., GROTON, COMM. 06249. J DRG CHEM 57 (6). 1992. 1814-1820. CCDEN: JOCEA Full Journal Title: Journal of Organic Chezistry Language: ENGLISH 7/3/23 (Item 2 from file: 55) 9138289 PIOSIS Number: 93123809 CORRECTION OF BR 92088484. CRYSTALLIZATION AND PRELIMINARY

DIFFRACTION DATA OF THE FAB FROSMENT OF A MONOCLONAL ANTIBODY

PROPERTY OF AUTUMN HOME COOM

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X-RAY

AGAINST

FITTE in black of all minimals. The behalf a summary of the parties of more about the minimal of the contract ALIAR. CRRATUM PUBLICHED IN FEBS FED EUR BICCHEM SOC LETT VOL. 292, 198, 1-2. 1991. P. 307 DEVAUX C; DEFENDINI M-L; ALIARI P M; ABERSEL C; GRANIER C; FONTECILLA-CAMPS J C LAB. BIOCHEM., FAC. MED.-MORD, BOULEVARD PIERRE DRAMARD, 13326 MARSEILLE CEDEX 15, FR. FERS (FED EUR BIOCHEM SOC) LETT 285 (1/2). 1991. 64-65. CODEN: FEBLA Full Journal Title: FEBS (Federation of European Biochemical Societies) Letters Language: ENGLISH 7/3/29 (Item 3 from file: 55) 8559708 BIOSIS Number: 92023708 A SANDWICH ENZYME-IMMUNCASSAY FOR DETECTING SNAME VENOM GE B; YU H; WANG J; XU Y SHANDKAI INST. BIGCKEMISTRY, ACABEMIA SINICA. ZCOL RES 12 (1). 1991. 79-83. CODEN: DOYAD Full Journal Title: Zoological Research Language: CHINESE 7/3/33 (Item 4 from file: 55) 7749725 BICSIS Number: 90117725 ISOLATION STRUCTURE ELUCIDATION AND SYNTHESIS OF MOVEL HYDROXYLAMINE-CONTAINING POLYAMINES FROM THE VENCY OF THE AGELENOPSIS-APERTA SPIDER JASYS V J; KELBAUGH P R; NASON D M; PHILLIPS D; ROSNACK X J; SACCOMANO N A; STROH J G; VOLICHANN R A CENTRAL RES., PFIZER INC., GROTCH, COMM. 06340. J AM CHEM SOC 112 (18), 1990, 6696-6704, CODEN: JACSA Full Journal Title: Journal of the American Chemical Society Language: EMSLISH (Item 5 from file: 55) 7/3/31 BIGGIS Number: 83038860 7110115 USING AFFINITY CHROMATOGRAPHY TO ISOLATE ALPHA LATROTOXIN FROM OF THE SPIDER LATRODECTUS-TREDECIMOUTTATUS DALIMOV E I; KREYMOV SH K; SALIKHOV SH I A.S. SADYKOV INST. BIGGRG. CHEM., ACAD. SCI. UZB. SSR, TASHKENT, USSR. KHIM PRIR SOEDIN (TASHK) 0 (5), 1983, 679-632, CODEN: KPSUA Full Journal Title: Khimiya Prirodnykh Spedinenii (Tashkent) Language: RUSSIAN 7/3/32 (Ites 6 from files 55) 7071659 BICSIS Number: 87132130

PURIFICATION OF FAR'-2 ANTI-SNAKE VENSA BY SAPRYLIC ASIG A FAST

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FOR OBTAINING IGG FRAGMENTS WITH HIGH NEUTRALIZATION ACTIVITY PURITY AND YIELD SANTOS M C D; LIMA M R D; FURTADO G C; COLLETTO G M D D; KIPNIS T L; SILVA W D D DEP. IMMUNOL., INST. CIENCIAS BIOMED., UNIV. SAO PAULO, BRAZIL.

TOXICON 27 (3). 1989. 297-304. CODEN: TOXIA Full Journal Title: Toxicon Language: ENGLISH

7/3/33 (Item 1 from file: 144)
10226038 PASCAL No.: 92-0431941
Analysis of a mastoparan P isolated from the hornet (Vespa Basalis) venom
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10226038 PASCAL No.: 92-0431941
Analysis of a mastoparan P isolated from the hornet (Vespi Basalis) venom
by fast atom bombardement mass spectrometry with B/E linked scan
YONG-CHIEN LING; BANG-JI LIU; JIRN-MEHNS LO; CHEW-LANG HO
National Tsing Hua univ., dep. chemistry, Hsinchu 300, Taiwan
Journal: Spectroscopy letters, 1992, 25 (2) 245-255
Language: English

7/3/34 (Item 2 from file: 144)
09555774 PASCAL No.: 91-0246201
Structure and biological activities of a new mastoparan isolated from the vencm of the hornet Vespa basalis
CHEWN-LANG HC; LING-LING HWANG
Acad. sinica, inst. biological chemistry, Taipei 10798, Taiwan Journal: Biochemical journal: (London. 1984), 1991, 274 (2) 453-456
Language: English

7/3/35 (Item 3 from file: 144)

@9114143 PASCAL No.: 90-0282524

Platelet glycoprotein IIb-IIIa protein antagonists from snake venoms:
evidence for a family of platelet-aggregation inhibitors

DENNIS M S; HENZEL W J; PITTI R N; LIPARI M T; NAPIER M A; DEISHER T A;
BUNTING S; LAZARUS R A

Genentech, Inc., dep. biomolecular chemistry, South San Francisco CA
94080, USA

Journal: Proceedings of the National Academy of Sciences of the United
States of America (1935), 1990, 87 (7) 2471-2475

Language: English

7/3/36 (Item 1 from file: 72) 8528645 EMBASE No: 92204527 Monoclonal antibodies to toxin II from the scorpion Androctonus australis

11\_\_1 \_ \_ . . .

neutralizing
capacities

Yahi N.; Davaux C.; Manquelle P.; Defendini M.-L.; Sranier C.
CNSS USR 1655, Laboratoire de Diochizie, Faculte de Medecine Secteur
Nord, Boulevard Pictre Drozord, 13226 Marseille Cedex 15 France
TOXICON (United Kingdon), 1932, 32/7 (723-731) CODEN:
TOXIA ISSN:
2041-0101 ADDNIS ORDER NUMBER: 0041010192200753
LANGUASES: English SUMMARY LANGUASES: English

7/3/37 (Item 2 from file: 72)
8343871 EMBASE No: 92082617
Neutralization of kinin-releasing enzymes from viperid venous by antivenon IgS fragments
Al-Joufi A.; Bailey G.S.; Reddi K.; Smith D.C.
Department of Chemistry and Biological Chemistry, University of Essex,
Colchester, Essex CO4 3SQ United Kingdom
TOXICON (United Kingdom), 1991, 29/12 (1509-1511) CODEN: TOXIA ISSN:
C041-0101 ADDNIS GREER NUMBER: 02410101920202037
LANGUAGES: English SUMMARY LANGUAGES: English

7/3/28 (Itez 3 from file: 72)
8319262 EMBASE No: 91342920
Emratur: Crystallization and preliminary X-ray diffraction data of the
Fab fragment of a concolonal antibody against aparin, a bec venor
neurotoxic (FEBS Letters (1991) 283 (64-66))

Devaux C.; Defendini M.-L.; Alzar P.M.; Abergel C.; Smanier C.; Fontecilla-Camps J.C. FEBS LETT. (Netherlands), 1991, 292/1-2 (307) CODEN: FEBLA ISSN: CO14-5793 ADDNIS CRDER NUMBER: 201457939101451L LANGUAGES: English

7/3/39 (Item 4 from file: 72)
6212121 EMBASE No: 07049774
Use of antibodies specific to defined regions of scorpion alpha-toxin to study its interaction with its receptor site on the sodium channel
Ayeb E.M.; Bahraoui E.M.; Granier C.; Rochat R.
INSERM U172 and CMRS UN 553, Laboratoire de Picchicie, Faculte de

Medecine Secteur Nord, 13225 Marseille Cedem 15 FRANCE BIOCHEMISTRY (USA) , 1985, 25/21 (6071-6672) CODEN: BICHA LANGUAGES: ENGLISH

7/7/AP //fer 1 form files 751)

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asine acid neuro-transmitter antagonists and calcium channel
blockers
Patent Assignee: (NATU-) NATURAL PROD SCI INC; (PFIZ ) PFIZER INC
Author (Inventor): ERCCOMANO N A; VOLKMANO R A
Patent Family:
                                 Week
   CC Marban
              Kind
                      Date
   EP 395357
               2.
                       201031
                                   9044
                                          (Basic)
   PT 93376
                                   9050
                А
                        901120
   AU 9854535
                A
                       961103
                                   9101
                       921229
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   CA 2015505
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   FI 9002139
                        901029
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   DD 293959
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                                   9229
   DD 298412
                         920219
                                   2237
   CS 9002157
                  82
Priority Data (CC No Date): US 345131 (890428)
Applications (CC,No,Date): CS 902157 (900428); EP 90304397
(900424); JP
   90115704 (900501); ZA 903229 (900427); DD 343270 (900805)
7/3/9
          (Item 9 from file: 155)
07478500 00385500
  Cleavage of innuneglobulins by socjeni protesse A, from
the venom of
Bothrops poojeni.
  Assakura MT; Mandelbaum FR
  Servico de Bicquipira, Instituto Butantan, Sac Faulo, Brasil.
 Texicon 1990, 28 (5) p734-6, ISSN 0041-0101 Journal Code:
WIT
 Languages: ENGLISH
  Document type: JOURNAL ARTICLE
7/3/10
           (Item 10 from file: 155)
07060610 89362610
  Issunctoxicotherapy: present status and future trends.
 Scherrmann JM; Terrien N; Untimberea M; Pierson P; Denis M;
  Insert U.26, Hepital Fernand Widal, Paris, France.
  J Toxicol Clim Toxicol 1989, 27 (1-2) p1-35, ISSN 0731-3910
Journal Code: KAN
 Languages: EMSLISM
  Document type: JOURNAL ARTICLE; REVIEW; REVIEW, TUTORIAL
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028442348 MPI Acc No: 92-329348/44

New poly-amine(s) and polypeptide(s) from spider vence - are

XRAM Occ No: 090-142975

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7/3/11
           (Item 11 from filer 155)
25939841 39291841
 Molecular recognition in the activation of human blood
ecagulation factor
 Chattopadhyay A; Fair DS
 Department of Biochesistry, University of Taxas Health
Conten, Tylan
75710.
 J Biol Chem Jul 5 1989, 264 (19) p11035-43, ISSN 2021-9258
Journal Code: HIV
 Contract/Grant No.: HL 39242
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
           (Item 12 from file: 155)
7/3/12
05965705 89257705
 Purification of F(ab)2 anti-snake venom by caprylic acid: a
for obtaining IgS fraggents with high neutralization activity,
purity and
yield.
 dos Santos MC; D'Isperio Liba MS; Funtado SC; Colletto SM;
Kipnis TL;
Dias da Silva W
 Fundação Ezequiel Dias, Universidade Federal de Minas Carais,
           1989, 27 (3) p297-303, ISSN 0041-0101
 Toxicon
                                                       Journal
Code: VWT
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
7/3/13
          (Item 13 from file: 155)
23869252 39171252
 Neutralization of lethal potency and inhibition of enzymatic
activity of
a phospholipase A2 neurotomin, protomin, by non-precipitating
antibodies
(Fab).
 Chouset V: Jiang MS: Radvanyi F: Cwnby C: Bon C
 Laboratoire des Venins, Institut Pasteur, Paris, France.
 FEBS Lett Feb 13 1989, 244 (1) p167-73, ISSN 0214-5793
Journal Code: EUM
 Languagess ENGLISH
 Document type: JOURNAL ARTICLE
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C&ECEC42 88170C42
                                  protects
                                               husan tuzer
 Decay-accelerating
                       factor
cells from
complement-mediated cytotoxicity in vitro.
 Cheung MM; Walter EI; Smith-Monsah WH; Ratnoff WD; Tykominski
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ML; Medof ME

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Department of Pediatrics, Memorial Sloan Kettering Cancer
Center, New
York 18821.
J Clin Invest Apr 1988, 8i (4) p1122-8, ISSN 8621-9738
Journal Code: HS7
Contract/Grant No.: CA-39326; AI-24226; AI-23598
Languages: ENGLISH
Document type: JOURNAL ARTICLE

7/3/15 (Item 15 from file: 155)
86358393 88603393
Block of synaptic transmission in insect CNS by toxins from
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Block of synaptic transmission in insect CNS by toxins from the venom of the wasp Megascolia flavifrons (Fab.).

Piek T; Hue B; Mony L; Nakajima T; Pelhate M; Yasuhara T
Department of Pharmacology, University of Amsterdam, The Netherlands.

Comp Biochem Physiol [C] 1987, 87 (2) p287-95, ISSN 0742-8413

Journal Code: DNX

Journal Code: DNX Languages: ENGLISH Document type: JOURNAL ARTICLE

7/3/16 (Item 16 from file: 155)
66089508 87663508
Immunotherapy in the poisoned patient. Overview of present applications
and future trends.
Sullivan JB Jr
Med Toxicol Jan-Feb 1986, 1 (1) p47-60, ISSN 0112-5966
Journal Code: MDT
Languages: ENGLISH

Document type: JOURNAL ARTICLE; REVIEW

7/3/17 (Item 17 from file: 155)
66035955 87609955
Defining the role of complement in experimental pemphigus vulgaris in mice.
Anhalt GJ; Till GO; Diaz LA; Labib RS; Patel HP; Eaglstein NF J Immunol Nov i 1986, 137 (9) p2835-40, ISSN 0622-1767
Journal Code: IFB
Contract/Grant No.: R23-AM32490; R23-AM32079; R01-AM32599; + Languages: ENGLISH
Document type: JOURNAL ARTICLE

7/3/18 (Item 18 from file: 155)
05649020 85265020
Role of complement and the Fc portion of immunoglobulin 6 in immunity to
Venezuelan equine encephalomyelitis virus infection with glycoprotein-specific monoclonal antibodies.

Hernicus and uncut ra att theme na J Virol Sep 1985, 55 (3) p594-600, ISSN 0022-538X Journal Code: KCV Languages: ENGLISH Document type: JOURNAL ARTICLE 7/3/19 (Item 19 from file: 155) 65534749 85156749

Modification endotexin-induced hasmodynamic and haematological changes in the rabbit by methylprednisolone, F(ab1)2 fragments and

resmarinic acid.

Bult H; Herman AG; Rampart M

Br J Pharmacol Feb 1985, 84 (2) p317-27, ISSN 0007-1188

Journal Code: B66 Languages: ENGLISH

Document type: JOURNAL ARTICLE

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**85416832 85826832** 

a bradykinin-like compound in the venom of Megascoliakinin, Megascolia

flavifrons Fab. (Hymenoptera: Scoliidae).

Piek T; Mantel P; Van Ginkel CJ

1984, 78 (2) p473-4, Comp Biochem Physiol [C]

0742-8413

Journal Code: DNX Languages: ENGLISH

Document type: JOURNAL ARTICLE

7/3/21 (Item 21 from file: 155)

64766845 82249845

Neurotoxin-specific

immunoglobulins

accelerate

dissociation of the neurotoxin-acetylcholine receptor complex. Boulain JC; Nenez A

Science Aug 20 1982, 217 (4561) p732-3, ISSN 0036-8075

Journal Code: UJ7 Languages: ENGLISH

Document type: JOURNAL ARTICLE

7/3/22 (Item 22 from file: 155)

81056391 64228391

Inhibition of biological activity of mouse beta-nerve growth factor by

monoclonal antibody.

Warren SL; Fanger M; Neet KE

Science Nov 21 1986, 210 (4472) p910-2, ISSN 0036-8075

Journal Code: UJ7

Contract/Grant No.: A110148; CA27915

Languages: ENGLISH

Document type: JOURNAL ARTICLE

7/3/23 (Item 23 from file: 155) A1254140 4121191A

Immunopharmacological approach to Foreszan shock. Magai H; Murimoto Y; Koda A Microbiol Izzunol 1920, 24 (7) p649-55, Journal Code: MX7 Languages: EMELICH Boougent type: JOURNAL ARTICLE 7/3/24 (14ez 24 froz filt: 155) 2405360E 00174695 Taxing retention: izaunological requirements for saintaining an easily degradable antigem in vivo. Tew JG; Mandel TE; Miller GA Aust J Exp Biol Med Sci Rug 1979, 57 (4) p401-14, 00C4-945X Journal Code: 9FW Languages: ENGLIGH Document type: JOURNAL ARTICLS 7/3/25 (Item 25 from file: 155) 03571494 79248494 Immunological studies on pancreatic phospholipase A2. Antigenie characterization of the NH2-terminal region. Maijer H; Maddens MJ; Dijkman R; Slotboom AJ; da Kaas GM J Biol Chem Dec 10 1978, 253 (23) p8564-9, ISSN 0021-9258 Journal Code: HIV Languages: EMSLISH Document type: JCURNAL ARTICLE 7/3/26 (Item 26 from file: 155) 02909946 76090940 nephrotonic nephritis. I. The role of Guin≥a-pig

polymorphonuclear leucceytes and the effect of antibody subplass and fraggents in the heterologous phase. Sispson IJ; Azos N; Evans DJ; Thosson NM; Paters DX Clin Exp Issunol May 1975, 19 (3) p499-511, ISSN 0309-9104 Journal Ocde: 007 Languages: ENSLISH Document type: JOURNAL ARTICLE 7/3/27 (Item 1 from file: 55) 9149315 DICEIS Number: 93134815 NOVEL QUATERNARY RYMONIUM SALT-CONTAINING POLYRMINES FROM THE ASSLENOPSIO APERTA FUNNEL-VED SPIDER JASYS V J; KELBAUSH P R; MACON D M; PHILLIPS D; ROSMACK K J; FORMAN J T; ENCOCHANG N A; STRON J G; VOLKMANN R A CENTRAL REGERROH DIVIGION, PRIZER INC., CROTCH, CORT. 05340. J GRG CHEM 57 (6). 1992. 1814-1820. CODEN: JOCEA Full Journal Title: Journal of Organic Chemistry Language: ENGLISH

complement and

7/3/23 (Item 2 from file: 55) BICSIS Number: 92122829 9138903 - CORRECTION OF BA 92038464. CRYSTALLIZATION AND PRELIMINARY X-39Y DIFFRACTION DATA OF THE FAB FRAGMENT OF A MONOGLONAL ANTIBODY APAMIN A BEE VENOM NEUROTOXIN. CORRECTION OF AUTHOR NAME FROM PEDRO M. ALIAG. GRANTUM PUBLISHED IN FEBS FED EUR BIOCHEM SOC LETT VOL. 292. 199. 1-2. 1991. P. 307 DEVAUX C: DEFENDINI M-L: ALZARI P M: ABERGEL C: GRANIER C: FONTECILLA-CAMPS J C LAB. BICCHEM., FAC. MED. -MORD, ECULEVARD PIERRE DRAMARD, 13326 MARSEILLE CEDEX 15, FR. FEBS (FED EUR BICCHEM ECC) LETT 285 (1/2). 1991. 64-65. CODEN: FEELA Full Journal Title: FEBS (Federation of European Bischemical Societies! Letters Language: ENGLISH

7/3/29 (Item 3 froz file: 55)
3558703 BIOSIS Nuzber: 92023703
A SANDWICH ENZYME-IMMUNDASSAY FOR DETECTING ENAME VENOM
SE 3; YU M; WANG J; XU Y
SMANSHAI INST. BICOMEMISTRY, ADADEMIA SINICA.
ZOOL RES 12 (1). 1991. 79-83. CODEN: DOYAD
Full Journal Title: Zoological Research
Language: CHINESE

7/3/30 (Item 4 from file: 55)
7749725 BIOSIS Number: 90117725
ISCLATION STRUCTURE ELUCIDATION AND SYNTHESIS OF MOVEL
MYDROXYLAMINE-CONTAINING POLYAMINES FROM THE VENOM OF THE
RSELENCASIC-APERTO EPIDER
JASYS V J; KELEAUGH P R; NASON D M; PHILLIPS D; ROSNACK K J;
SACCOMAND N
A; STRCH J G; VOLKMANN R A
CENTRAL RES., PFIZER INC., GROTON, CONN. 06340.
J CM CHEM SCO 112 (18). 1990. 6696-6704. CODEN: JACSA
Full Journal Title: Journal of the Aperican Chesical Speiety
Language: ENGLISH

711611E BIGGIS NUMBER: 20028860

USING ACTINITY CHROMATOGRAPHY TO ISCLATE ALPKA LATACTOXIN FROM THE VENOM
OF THE SFIDER LATACOSCIUS-TREDECIMOUTTATUS

DALIMON 2 Z; MASYMEN CM N; SALIKUON CH I
A.S. SREYKON INST. BICCAS. CHEM., ACAD. SCI. UZB. SSR,
TAGISKOTT, USER.

(Itea 5 from file: 55)

7/3/31

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Full Journal Title: Whisiya Privatelih Cosdinarii (Tashkani)
 Languages RUCGIAN
7/3/32
           (Item & from file: EE)
7071659
          BICCIC Number: 07132180
 PUBLIFICATION OF FAR'-2 ANTI-CHARGE VEHICH BY CAPAVILIC ACID A FAST
FOR DATAINING IGS FRACMENTS WITH HIGH NEUTRALIZATION ACTIVITY
DURITY AND
YIELD.
 SANTOS M C D: LIMO M R D: FURTADO S C: COLLETTO S M D D: KIPNIS
SILVA U D D
 PEP. INMUNEL, INST. CIENCIAS RIGNED., UNIV. SAD PAULO, ERAZIL.
 TOXICON 27 (3), 1989, 297-304, CODEN: TOXIA
 Full Journal Title: Toxicon
 Language: ENGLISH
           (Item 1 from file: 144)
7/3/33
 Analysis of a pastoparan B isolated from the hornet (Vesps
Basalis) venco
by fast atom bombardement cass spectrometry with B/S linked scan
 YONG-CHIEN LING: BANG-JI LIU: JIRM-MEKNG LO: CHEW-LANG KO
 National Tsing Hua univ., dep. chemistry, Hsinchu 200, Taiwan
 Journal: Spectroscopy letters, 1992, 25 (2) 245-255
 Language: English
         (Itaz 2 fraz file: 144)
 7/3/34
 09555774 PASCAL No.: 91-0345201
 Structure and biological activities of a new pastoparan
isolated from the
venco of the hornet Vespa basalis
 CKEWN-LANG KO: LING-LING KHANG
 Acad. sinica, inst. biological chapistry, Taipai 10798, Taiwan
 Journal: Biochepical journal: (London, 1984), 1991, 274 (2)
453-456
 Language: English
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7/3/35 (Item 3 from file: 144)

00114143 PREDAL No.: 90-0282524

Platelet glycoprotein III-IIIa protein antagonisto from snake venous:
evidence for a family of platelet-aggregation inhibitors

DERMIS M E; KENZEL W J; PITTI N M; LIPARI M T; NAPIER M A;
DEISMER T A;
BUNTING G; LAZARUG R A

Genentech, Inc., dap. bicoolecular checistry, South San Francisco CA
94262, USA

Journal: Droceedings of the National Academy of Epienous of the United
Distance of Common Stocks of Section 22 (2) 9434 9435

which is the see that and the set in the mit-Language: English 7/3/33 - (Itas i fysa files 72) 8520345 EMERSE No: 92224527 Monoclonal antibodies to towin II from the scorpion Androctomus Mactor: Further characterization of cpitope specificities and nautralizing papapities Yahi M.; Devaux C.; Mansuelle P.; Defondini M.-L.; Smanier C. CMMS UMA 1455, Laboratoire de Bischizie, Faculte de Nord, Boulevard Pierre Drazard, 13325 Marseille Cadex 15 France TOXICON (United Kingdom) , 1992, 20/7 (722-731) CODEN: TOXIA ISSN: 0041-0101 ADDNIS CRDER NUMBER: 0041010192000753 LANGUAGES: English SUMMARY LANGUAGES: English 7/3/37 (Item 2 from file: 72) 0342871 EMZASE No: 92020517 Neutralization of kinin-releasing enzypes from viperid venous by antivenos IgS fragments Al-Joufi A.; Bailey G.S.; Reddi M.; Saith D.C. Department of Chemistry and Biological Chamistry, University Colchester, Essax CC4 380 United Kingdoo TOXICON (United Kingdom), 1991, 29/12 (1509-1511) CODEN: TOXIA ISSN: 2241-0101 ADDNIS ORDER NUMBER: 0241010192002097

LAMBURSES: Emplish SUMMARY LAMBURSES: Emplish

7/2/39 (Item 3 from file: 72)
8310262 EMBASE No: 91340920
Erratum: Crystallization and preliminary X-ray diffraction data of the
Fab fragment of a conscional antibody against apacin, a bas venomentowin (FEDS Letters (1991) 285 (54-65))
Devaum C.; Defending M.-L.; Alzar P.M.; Abargel C.; Smanier C.;
Contemilla-Camps J.C.
FEDS LETT. (Netherlands) , 1991, 292/1-2 (307) CODEN: FEDLO ISSN:
C214-5792 ADCMIS CAMER NUMBER: C014579391014511
LONGUAGES: English
7/3/39 (Item 4 from file: 72)

13:3121 EMBRCE No: 07040774

"Use of antibodies specific to defined regions of scomplon alpha-towin to study its interaction with its receptor site on the sodium channel.

"The Park of the Common Definite Park of the sodium channel."

INSERM U172 and CNRS UA 553, Laboratoire de Biochimie, Faculte de
Medecine Secteur Nord, 13326 Marseille Cedex 15 FRANCE
BIOCHEMISTRY (USA) , 1986, 25/21 (6671-6678) CODEN: BICHA
LANGUAGES: ENGLISH

7/3/40 (Item 1 from file: 351) 668442348 WPI Acc No: 98-329348/44

XRAM Acc No: C90-142975

New poly-amine(s) and polypeptide(s) from spider venom - are eccitatory

amino acid neuro-transmitter antagonists and calcium channel blockers

Patent Assignee: (MATU-) NATURAL PROD SCI INC; (PFIZ ) PFIZER INC

Author (Inventor): SACCOMANC N A; VOLKMANN R A Patent Family:

CC Number Kind Date Week 901031 EP 395357 Α 9844 (Basic) PT 93876 901120 9050 901108 9101 AU 9054535 Α A 901029 9101 NO 9001881 Α CA 2015505 901028 9104 FI 9882139 Α 901029 9107 Α 910123 9116 JP 3014551 HU T55415 T 910528 9127 Α 901121 9131 CM 1047078

ZA 9003229 Α 911224 9205 916919 9208 DD 293959 A DD 298412 A5 920220 9229 A2 CS 9662157 920219 9237

Priority Data (CC No Date): US 346181 (596428)

Applications (CC,Ne,Date): CS 902157 (900428); EP 90304397 (900424); JP 90115704 (900501); ZA 903229 (900427); DD 343270 (900806) ?ds

Set Items Description S1 591 ANTIVENIN 52 SI AND ( F(AB) OR FAB FRAGMENT OR F(AB)2) **S**3 2 SI AND ( F(AB) OR FAB(W) FRAGMENT OR FAB OR FAB2 ) 54 23862 VENOM **S**5 19302 FAB OR FAB2 **S6** 72 54 AND 55 **S7** 40 RD (unique items) ?s s5 and polyacrylamide

19382 S5 146486 POLYACRYLAMIDE S8 586 S5 AND POLYACRYLAMIDE ? s s8 and (pepsin or papain) 586 S8 15002 PEPSIN 10290 PAPAIN 92 S8 AND (PEPSIN OR PAPAIN)

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59

)))Duplicate detection is not supported for File 356.
)))Duplicate detection is not supported for File 351.

)))Records from unsupported files will be retained in the RD set.

...examined 50 records (50)
...completed examining records
S10 75 RD (unique items)
?t s10/3/1-75

10/3/1 (Item 1 from file: 155)

**98177636** 92315636

Thiol groups and other chemical characteristics of rat monoclonal

issunoglobulin A.

Gertler S; Young NM

Institute for Biological Sciences, National Research Council of Canada,

Ottawa, Ontario.

**07849323** 

Dose-dependent

poisoning by goat

91368323

colchicine-specific Fab fragments.

Comp Biochem Physiol [B] (ENGLAND) Jun 1992, 102 (2) p377-81, ISSN

6365-6491 Journal Code: DNV

Languages: ENGLISH

Document type: JOURNAL ARTICLE

10/3/2 (Item 2 from file: 155) 08080766 92218766 Single-step purification of F(ab1)2 fragments of mouse antibodies (immunoglobulins G1) by hydrophobic interaction high performance liquid chromatography using TSKgel Phenyl-5PW. Morimoto K; Inouye K Biotechnology Research Laboratories, TOSOH Corporation, Kanagawa, Japan. J Biochem Biophys Methods (NETHERLANDS) Mar 1992, 24 (1-2) p107-17, ISSN 6165-622X Journal Code: H94 Languages: ENGLISH Document type: JOURNAL ARTICLE (Item 3 from file: 155) 16/3/3

reversal of

murine

acute

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Institut National de la Sante et de la Recherche Medicale 125, Mopital
Fernand Widal, Paris, Francs.
 Codo: WMD
 Languages: CNGLICH
 Document type: JOURNAL ARTICLE
          (Item 4 from file: 155)
10/2/4
07823692 91342692
 The Fab/c fragment of IgG produced by cleavage at cyanocysteine
residues.
 Wines ED; Easterbrook-Seith SB
 Department of Biochemistry, University of Sydney, M.S.M.,
 Mol Torunol Rug 1991, 28 (8) p855-63, ISSN 0161-5090
Journal Codes
XE1
 Languages: EMSLISH
 Document type: JOURNAL ARTICLE
10/3/5
         - (Idea 5 frac file: 155)
07821660 91340600
 Vascantive intestinal paptide hydrolysis by antibody light
chains.
 Mei S; Mady B; Eliland EM; Paul S
 Department of Therespology, University of Rebraska Medical
Center, Omaha
68199-6200.
 J Diel Chae | Aug 25 1991, | 265 (24) | p15571-4, | ISSN 2021-9258
Journal Code: KIV
 Contract/Brant No.: ML40348; KL44125; KL22217
 Languages: ENGLISH
 Decadent type: JCURNAL ARTICLE
10/3/6 (Itac & from file: 155)
C7591E47 91210547
 Proteclytic digestion of cause IgE.
 Maba St Misonoff A
 Department of Biology, Brandeis University, Walthau, MA 02254.
 J Issued Mathods Apr 8 1991, 138 (1) p15-22, ISSN
0822-1759
Journal Octor ISE
 Contract/Grant No.: AI-22068
 Languages: ENSLISH
 Domument type: JOURNAL ARTICLE
 10/2/7
          - 115to 7 from Filte 155)
27571709 91092709
 The specificity of the IgO receptor purified from human
meutrophils.
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manufacture (); as character as is as amagine go (i) amount of seattle () (by

Department of Pathology, University of Dundee, Ninewell's Hospital Medical
School, Scotland, U.K.
Biochem J Nov 15 1990, 272 (1) p159-65, ISSN 6264-6621
Journal Code: 9YO
Languages: ENGLISH
Document type: JOURNAL ARTICLE

16/3/8 (Item 8 from file: 155)
67569635 91688635
Preparation and crystallization of a human immunodeficiency

07569635 91088635

Preparation and crystallization of a human immunodeficiency virus p24-Fab complex.

Prongay AJ; Smith TJ; Rossmann MG; Ehrlich LS; Carter CA; McClure J

Department of Biological Sciences, Purdue University, West Lafayette, IN

Lafayette, IN 479**8**7.

Prec Natl Acad Sci U S A Dec 1990, 87 (24) p9980-4, · ISSN 0027-8424

Journal Code: PV3

Contract/Grant No.: AI25993; AI27316

Languages: ENGLISH

Document type: JOURNAL ARTICLE

10/3/9 (Item 9 from file: 155) 67552858 91071858

Purification and characterization of an outer membrane protein adhesin

from Haemophilus parainfluenzae HP-28.

Lai CH; Bloomquist C; Liljemark WF

Department of Diagnostic School of Dentistry, University of

Minnesota,
Minneapolis 55455.
Infect Immun Dec 1990, 58 (12) p3833-9, ISSN 0619-9567
Journal Code: 607
Contract/Grant No.: R37-DE04614
Languages: ENGLISH
Document type: JOURNAL ARTICLE

40/3/10 (Item 10 from file: 155) 87400795 90307795

Characterization of a maternal type VI collagen in Xenopus embryos

suggests a role for collagen in gastrulation.

Otte AP; Roy D; Siemerink N; Koster CH; Hochstenbach F; Timmermans A;

Durston AJ

Hubrecht Laboratory, Netherlands Institute for Developmental Biology, Utrecht.

J Cell Biol Jul 1990, 111 (1) p271-8, ISSN 0621-9525 Journal Code:

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Languages: ENGLISH
 Document type: JOURNAL ARTICLE
 10/3/11
             (Item il from file: 155)
67349288
          98256288
 A proteclytic enzyme secreted
                                            Proteus
                                                     #irabilis
immuneglobulins of the immunoglobulin A1 (IgA1), IgA2, and Ig6
isetypes.
 Loomes LM; Senior BW; Kerr WA
 Department of Pathology, Dundee University Medical School,
Ninewells
Hospital, Scotland.
                Jun 1990, 58 (6) p1979-85, ISSN 6019-9567
  Infect Immun
Journal Code: GO7
 Languages: ENGLISH
 Document type: JOURNAL ARTICLE
 10/3/12
             (Item 12 from file: 155)
67345988
          96252988
 Adaptation of fluorescence polarization
                                                immunoassay
the assay of
macromolecules.
 Urios P; Cittanova N
 Departement
                       Biochimie,
                                     U.F.R.
                                               Biomedicale des
                de
Saints-Peres, Paris,
France.
  Anal Biochem Mar 1990, 185 (2) p308-12, ISSN 0603-2697
Journal Code: 4NK
 Languages: ENGLISH
  Document type: JOURNAL ARTICLE
 10/3/13
             (Item 13 from file: 155)
07062848 89364848
                   studies
                              of a murine polyreactive Ig62b
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J Biochem (Tokyo) May 1989, 165 (5) p863-12, ISSN 6621-924X

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Matsumura S; Ohmori K; Chiba T; Kumon A

Department of Biochemistry, Saga Medical School.

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Dias da Silva W

Fundacao Ezequiel Dias, Universidade Federal de Minas Gerais, Brazil.

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Laboratory of Pathology, Aichi Cancer Center Research Institute, Nagoya, Japan.

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Department of Immunology, Fundacion Jimenez Diaz, Madrid, Spain.

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Bowles M; Johnston SC; Schoof DD; Pentel PR; Pond SM

Department of Medicine, University of Queensland, Princess

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Erntell M; Myhre EB; Sjobring U; Bjorck L

University of Lund, Department of Infectious Diseases, University

Hospital, Sweden.

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Department of Pediatrics, Yamaguchi University School of Medicine.

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 Document type: JOURNAL ARTICLE
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Lykkegard S

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serum IgA)
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5286014 BIOSIS Number: 81053321
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ANTIGEN-BINDING IMMUNOGLOBULIN A FRAGMENTS FROM PAPILLARY DERMIS OF
UNINVOLVED SKIN BY PEPTIDE DIGESTION
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ARCH DERMATOL REG 278 (1). 1985. 44-48. CCDEN: ADRED Fall Journal Title: Archives of Dermatological Research

Language: ENGLISH

10/3/74 (Item & from file: EE) 4885648 BIOSIS Number: 80013359 DERMATITIS HERPETIFORMIS BIOCHEMICAL PROPERTIES OF THE GRANULAR DEPOSITS OF IMMUNOGLOBULIN A IN PAPILLARY DERMIS CHARACTERIZATION OF SODIUM DODECYL SULFATE-SCLUBLE IMMUNOSLOBULIN A-LIKE MATERIAL AND POTENTIALLY ANTIGEN-BINDING IMMENOGLOBULIN & FRAGMENTS RELEASED BY PEPSIN EGELRUD T; BACK C DEPARTMENT OF DERMATOLOGY, UNIVERSITY HOSPITAL, S-921 85 LIMEA, SWEDEN. J INVEST DERMATCL 84 (4). 1985. 239-245. CODEN: JIDEA Full Journal Title: Journal of Investigative Dermatology Language: ENGLISH

10/2/75 (Item 7 from file: 55)
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BJORCK L; KRONVALL 6
DEP. MED. MICROBIOL., SOLVEGATAN 23, S-223 62 LUND, SWED.
J IMPLINOL 133 (2). 1984. 969-974. CODEN: JOIMA
Full Journal Title: Journal of Impunology
Language: ENGLISH

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1. 5,180,826, Jan. 19, 1993, Brain-derived neurotrophic factor; Yves-Alain Barde, et al., 536/23.51; 435/69.1, 69.3, 172.3, 320.1;

530/399, 412 [IMAGE AVAILABLE]

US PAT NO: 5,180,820 (IMAGE AVAILABLE)

L3: 1 of

39

ABSTRACT:

The present invention relates to nucleic acid sequences encoding brain derived neurotrophic factor (BDNF), as well as BDNF protein

derived neurotrophic factor (BDNF), as well as BDNF protein produced in

ту не в од межници болени поличини моници оборителници да веня на подаваления ā.... derivatives thereof. In addition, the invention related to pharascologic compositions and thereposite uses of EDMF, having provided, for time, the ability to generate sufficient quantities of substantially pure BDAT for plinical use. The invention also relates to antitodies directed toward BENF on fragments thereof, having provided a method for gamayating sufficient issumagen. Further, by percitting a comparison of the acid sequences of BOMF and NGF, the present invention provides for the identification of homologous regions of number amid sequence between BDNF and NGF, theraby defining a BDNF/NGF gene family; the invention provides a sethed for identifying an displating additional

machers of
this gene family.
2. 5,175,273, Dec. 29, 1992, Nucleic acid intercalating agents;
Norbert
M. Bischofberger, et al., 525/27, 1.1, 18.7, 22, 23, 29, 29

US PAT NO: 5,175,273 [IMAGE AVAILABLE] L3: 2 of 39

# ABSTRACT:

CIMAGE AVAILABLEI

Pyridinone or pyrisidinone nucleoside bases containing fused arosatic polycyclic rings are provided. These polycyclic nucleosides are incorporated into oligonucleotides and hybridized to complementary nucleic acid. Fluorescence spectroscopy and thermal denaturation profiles provided evidence that the polycyclic base is intercalated into the resulting duplem. The fused polycyclic ring systems optionally are substituted with reactive species which inactivate complementary nucleic acids. The aligonucleotides of this invention are useful as improved probas, diagrostic reagents, or for alsaving or derivatizing prodoteroined descins within nucleic acids.

2. 5,171,864, Dec. 15, 1992, Di-45-hydron,-2(5D)-2-cke-4-funyl)mothylfuryl)methyl-alpha,omega-dialkanoic acid amides as anti-inflammatory agents; Gary C. M. Lee, 549/222, 318 [IMAGE AVAILABLE]

US PAT NO: 5,171,864 [IMAGE AVAILABLE] L3: 3 of 39

# ABSTRACT:

Compounds of the formula ##STR1## in which R. sub.1 independently is H or alkyl of 1 to 20 carbons, CO-R. sub.2, CO-O-R. sub.2, CO-NH-R. sub.2, or PO(OR. sub.2). sub.2 or PO(OR. sub.2)R. sub.2, where R. sub.2 independently is H, alkyl of 1 to 20 carbons, phenyl, or lower alkyl substituted phenyl or halogen substituted phenyl; A is (CH. sub.2). sub.n where n ranges between 5 to 36, or A is a a divalent branch chained alkyl radical, or cycloalkyl radical, having a total of 5 to 30 carbons, and X is O or NH, have anti-inflammatory activity.

4. 5, 169,963, Dec. 8, 1992, Di-(5-hydroxy-2(5H)2-oxo-4-furyl)alkylmethylalpha,omega alkanedioates and N,N-bis-(5-hydroxy-2(5H)2-oxo-4-furyl)alkylmethyl-alpha,omega-dialkanoic acid amides as anti-inflammatory agents; 6ary C. M. Lee, 549/222, 318 [IMAGE AVAILABLE]

US PAT NO: 5,159,963 [IMAGE AVAILABLE] L3: 4 of

39

### ABSTRACT:

Compounds of the formula ##STRI## in which R. sub.1 independently is H or alkyl of 1 to 29 carbons, CO-R. sub.2, CO-O-R. sub.2, CO-MH-R. sub.2, or PC(OR. sub.2). sub.2 or PC(OR. sub.2). sub.2, where R. sub.2 independently is H, alkyl of 1 to 20 carbons, phenyl, or lower alkyl substituted phenyl or halogen substituted phenyl; A is (CH. sub.2). sub.n where n ranges between 8 to 30, or A is a a divalent branch chained alkyl radical, or cycloalkyl radical, having a total of 3 to 30 carbons; R. sub.3 independently is an alkyl group having 4 to 20 carbons, and X is C or NH, have anti-inflammatory activity.

5. 5,169,933, Dec. 8, 1992, Covalently-linked complexes and

enhanced cytotoxicity and imaging; David C. Anderson, et al., 536/391.3; 424/1.1, 9, 85.91; 536/366, 367, 368, 313, 323, 324, 325, 326, 327, 328, 329, 330, 351, 377, 391.1, 391.5, 391.7, 391.9, 395, 399, 403, 468, 409, 416 [IMAGE AVAILABLE]

US PAT NO: 5,169,933 [IMAGE AVAILABLE] L3: 5 of 39

# ABSTRACT:

are also described.

Covalently-linked complexes (CLCs) for targeting a defined population of cells, comprising a targeting protein; a cytotoxic agent; and an enhancing moiety, wherein the enhancing moiety is capable of promoting CLC-target cell interaction are disclosed. Methods for using the claimed CLCs to obtain enhanced in vivo cytotoxicity and enhanced in vivo imaging

6. 5,162,504, Nov. 10, 1992, Monoclonal antibodies to a new antigenic marker in epithelial prostatic cells and serum of prostatic cancer patients; Julius S. Heroszewicz, 530/388.2; 435/7.23, 70.21, 240.27; 530/388.8 [IMAGE AVAILABLE]

US PAT NO: 5, 162, 584 [IMAGE AVAILABLE] L3: 5 of

39

### ABSTRACT:

Monoclonal antibodies to prostatic cells, are produced by a hybridoma

formed by fusing mouse lymphocytes and mouse myeloma cells. The monoclonal antibodies show specificity for a non-soluble, membrane

associated, organ specific antigenic determinant limited in its distribution to normal and neoplastic, human prostate epithelial cells.

The monoclonal antibodies, specifically 7E11-C5 monoclonal antibodies,

may be suitable for diagnostic uses.

7. 5,159,685, Oct. 27, 1992, 2-anilino phenylacetic acid derivatives as inhibitors of PLA.sub.2 and lipoxygenase; Amedec A. Failli, et al., 548/318.1, 341.1, 342.5 [IMAGE AVAILABLE]

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# ABSTRACT:

There are disclosed compounds of the formula ##STRI## wherein R is

hydroxy, lower alkoxy or lower alkoxyamino;

R. sup. 1 is hydrogen or A(CH. sub. 2). sub. n 0--;

R.sup.2 is hydrogen or A(CH.sub.2).sub.n 0--, with the proviso that one

of R. sup. 1 and R. sup. 2 is hydrogen;

n is 1-2;

ř

A is phenoxyethyl, phenoxyphenyl or a group having the formula ##STR2##

R.sup.3 is hydrogen, lower alkyl or phenyl; R.sup.4 is hydrogen or

lower alkyl; or

R. sup. 3 and R. sup. 4 taken together form a benzene ring;

R. sup. 5 is hydrogen or lower alkyl;

R. sup. 6 is hydrogen, halo or lower alkyl;

and the pharmacologically acceptable salts thereof, and their use in the

treatment of inflammatory conditions, such as rheumatoid arthritis,

ulcerative colitis, psoriasis and other immediate hypersensitivity

reactions; in the treatment of leukotriene-mediated naso-bronchial

obstructive air-passageway conditions, such as allergic rhinitis,

allergic bronchial asthma and the like; and as gastic cytoprotective agents.

8. 5,156,846, Oct. 20, 1992, Amine-containing porphyrin derivatives;
John W. F. Goers, et al., 424/85.91; 514/410 [IMAGE AVAILABLE]

US PAT NO: 5,156,840 [IMAGE AVAILABLE] L3: 8 of 39

### ABSTRACT:

The invention relates to amine-containing porphyrin derivatives. Theporphyrins can be used as photosensitizers which are useful as

therapeutic agents. Also described are methods for preparing conjugates in which a porphyrin derivative is covalently attached to an antibody or antibody fragment. In vivo therapeutic methods utilizing the conjugates

are also desired.

9. 5,135,736, Aug. 4, 1992, Covalently-linked complexes and

enhanced cytotoxicity and imaging; David C. Anderson, et al.,
424/1.1, 9,
85.91, 94.3; 435/188; 514/8, 12, 21; 536/387.1, 388.15, 388.8,
389.7,
391.3, 391.7, 395 [IMAGE AVAILABLE]

US PAT NO: 5,135,736 [IMAGE AVAILABLE] L3: 9 of 39

# ABSTRACT:

Covalently-linked complexes (CLCs) for targeting a defined population of cells, comprising a targeting protein or peptide; a cytotoxic agent; and an enhancing moiety, wherein the enhancing moiety is capable of promoting CLC-membrane interaction are disclosed. Methods for using the claimed CLCs to obtain enhanced in vivo cytotoxicity and enhanced in vivo imaging are also described.

5,120,537, Jun. 9, 1992, Factor Xa based anticoagulant compositions;
 Charles T. Esmon, et al., 424/94.64; 435/69.6, 212, 226; 514/2, 12, 21;
 530/381 [IMAGE AVAILABLE]

US PAT NO: 5,120,537 [IMAGE AVAILABLE] L3: 10 of

ABSTRACT:

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An anticoagulant composition containing an effective amount of factor Xa having the active serime site inactivated that functions rapidly effectively in vivo to suppress coagulation. In a preferred embodiment, Factor Xa, a serine esterase that forms a complex with Factor Va, and phospholipid to catalyze prothrosbin activation, is first inactivated with an active site inhibitor, such as dansyl-glugly-arg-chloromethyl ketone, to form inactivated factor Xa. In another embodiment, is expressed from a gene sequence wherein the portion encoding serine region is modified. The inactivated protein retains the ability to bind to endogenous factor Va in vivo, and has a half-life of approximately ten hours. Administration of inactive factor Xa to

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Xa-Va

complexes in vivo, thereby inhibiting coagulation.

11. 5,112,739, May 18, 1992, Enzyme controlled release system; Frank A.

Meneghini, et al., 435/14, 4, 18, 19, 21, 24, 183; 436/546 [IMAGE

# AVAILABLE)

US PAT NO: 5,112,739 [IMAGE AVAILABLE] L3: 11 of

# ABSTRACT:

This invention relates to an enzyme-controlled release method for the

release of a leaving group comprising:

contacting a compound represented by the formula ##STR1## wherein R;

R.sub.1, R.sub.2 and R.sub.3 each independently is hydrogen, a substituent affecting the mobility or reactivity of the compound or a

substituent including a biologically active group;

X is leaving group;

Z is an enzyme substrate moiety;

--CR.sub.2 R.sub.3 X is either ortho or para to the --O--Z moiety;

with an active capable of cleaving said enzyme substrate moiety  $\mathcal I$  from

said compound;

whereby said leaving group X is released from said compound.

12. 5,**086,00**2, Feb. 4, 1992, Erythrocyte agglutination assay;

Carmel J.
Hillyard, et al., 436/540; 422/61; 435/7.25; 436/501, 519; 530/387.3, 388.7, 389.1, 866 [IMAGE AVAILABLE]

US PAT NO: 5,086,002 [[MAGE AVAILABLE] L3: 12 cf

# ABSTRACT:

In a novel, erythrocyte agglutination assay, the agglutination reagent

comprises at least one erythrocyte binding molecule coupled to at least

one specific analyte binding molecule wherein the erythrocyte binding

molecule does not cause agglutination when incubated with erythrocytes in

the absence of analyte (in the case of a direct assay) or analyte binding

reagent (in the case of an indirect assay). Preferably, the erythrocytes

sample is assayed. Mixtures of conjugates and conjugates of analogues with erythrocyte binding molecules may also be used as agglutination reagents. The reagents and their use in direct or indirect assays is disclosed.

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13. . 5,681,261, Jan. 14, 1992, 4-(1-hydroxy-2-N-substituted sulfonamida) ethyl-5-hydroxy-2(5H)-furanones and 4-(N-substituted sulfonamido)-2ethenyl-5-hydroxy-2(5H)-furanones as anti-inflammatory agents; Lee, 549/222; 544/229, 337, 379, 383; 549/214, 313, 318 [IMAGE AVAILABLE]

5,081,261 [IMAGE AVAILABLE] L3: 13 of US PAT NO: 39

# ABSTRACT:

Compounds of Formula 1, and of Formula 2, ##STRI## in which R. sub. 1 is H CO--R. sub. 1. sup. ₹ alkyl to 26 carbons, CO--O--R. sub. i. sup. \* CO--NH--R. sub. 1. sup. \* or PO(OR. sub. 1. sup. \*). sub. 2 or PO(OR. sub.1. sup. \*) R. sub.1. sup. \* where R. sub.1. sup. \* independently alkyl of 1 to 20 carbons, phenyl, or substituted phenyl; R.sub.2 is H. alkyl of 1 to 20 carbons, or R. sub. 2 and Y jointly represent a heterocycle which incorporates the sulfonamide mitrogen in the

ring as a heteroatom: R.sub.3 is H or alkyl of 1 to 20 carbons: X is H. R. sub. 4, CO---O---R. sub. 4, CO--NH--R. sub. 4, CO--R. sub. 4, CO--N--(R. sub. 4). sub. 2, PO(OR. sub. 4), sub. 2 er PO(OR. sub. 4) R. sub. 4, and independently is H, phenyl, substituted phenyl, alkyl of 1 to 20 carbons or is to 26 carbons substituted with a hydroxyl, alkoxy, substituted amino, thioalkoxy, with a G--COR.sub.4.sup.\* group or with a COR. sub. 4. sup. \* group where R. sub. 4. sup. \* is H, lawer alkyl, OH, OR. sub. 4. sup. ^\_ NH. sub. 2, NHR. sub. 4. sup. \_\_\_ or N(R. sub. 4. sup. \_\_\_). sub. 2 group where R. sub. 4. sup. \_\_ independently is H or lower alkyl, with the provise that when X is CO--O--R. sub. 4 or is CO--NH--R. sub. 4 then R. sub. 4 is not

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to 50.

carbons, or is alkyl of 1 to 26 carbons substituted with a hydroxyl,

alkoxy, substituted amino, thicalkoxy, 0--PO(OR.sub.5).sub.2, 0--PO(OR.sub.5)R.sub.5, 0--SO.sub.3 H, 0--SO.sub.2 R.sub.5, 0--COR.sub.5,

or COR.sub.5 group where R.sub.5 is H, lower alkyl, OH, OR.sub.5.sup.\*,

NH. sub. 2, NHR. sub. 5. sup. \* or N(R. sub. 5. sup. \*). sub. 2 group where R. sub. 5. sup. \* is lower alkyl, or R. sub. 2 and Y jointly represent a

heterocycle which incorporates the sulfonamide nitrogen in the ring as a

heteroatom, with the provise that when Y is an alkyl substituted with

O--PG(OR.sub.5).sub.2 or with O--PO(OR.sub. 5)R.sub.5 then R.sub.5 is not

OH, are disclosed. The compounds possess anti-inflammatory activity.

14. 5,071,935, Dec. 10, 1991, Ribonucleotide reductase inhibitors; Yvan Guindon, et al., 514/13, 15, 17, 18, 19; 530/326, 327, 328, 329 [IMAGE AVAILABLE]

US PAT NO: 5, 671, 835 [IMAGE AVAILABLE] L3: 14 of 39

## ABSTRACT:

Disclosed herein are peptides of the formula

Y--R. sup. 1 --R. sup. 2 --R. sup. 3 --R. sup. 4 --R. sup. 5 --R. sup. 6 --Z

wherein R.sup.1 to R.sup.5 are designated amino acid residues; R.sup.6 is

Phe, homoPhe or an amino acid residue derived from 2-amino-3cyclohexylpropionic acid, 2-amino-3-(4-(lower alkoxy)phenyl) propionic

acid or 2-amino-3-(4-halophenyl)propionic acid; Y is Phe, desamino-Phe.

(lower alkanoyl)-Phe, p-haloPhe, Tyr, desamino-Tyr or (lower alkanoyl)-Tyr, or Y is the decapeptide radical W--Val--R.sup.7
--Ser--R.sup.8 --R.sup.9 --Thr--Glu--R.sup.10 --Ser--Phe wherein W is

hydrogen or lower alkanoyl and R.sup.7 to R.sup.16 are designated amino

acids residues, or Y is a fragment of the decapeptide radical wherein

from one to nine of the amino acid residues (i.e. Val to Ser) may be

deleted serially from the amino terminus of the decapeptide radical; and

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peptides inhibit mammalian ribonucleotide reductase and are preventing or ameliorating abnormal cell proliferation.

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1991, 5,866,789, Nov. 19, Targeting substance-diagnostic/therapeutic agent conjugates having Schiff base linkages; Ananthachari Srinivasan, et al., 530/391.5, 363, 391.9 [IMAGE AVAILABLE]

L3: 15 of US PAT NO: 5,066,789 [IMAGE AVAILABLE] 39

# ABSTRACT:

substance-diagnostic/therapeutic agent conjugates Targeting joined by stabilized Schiff base or hydrazone linkages are disclosed. In slow release carrier-drug pharmaceuticals are described. The diagnostic and therapeutic conjugates and pharmaceuticals of the present provide certain advantages relating to in vivo administration, including controlled release of the active agent at a target site.

5,666,671, Nov. 19, 1991, Ellagic acid derivatives as phospholipase A. sub.2 inhibitors; Craig E. Caufield, 514/453 [IMAGE AVAILABLE]

5,066,671 (IMAGE AVAILABLE) US PAT NO: L3: 16 of

39

n=1-3;

There are disclosed a method for the treatment or prevention of immunoinflammatory conditions by administering to a mammal an amount of a compound having the formula: ##STR1## wherein R. sup. 1. R. sup. 2, R. sup. 3, and R. sup. 4 are each, independently, hydrogen, alkyl, aralkyl, aryl, or ##STR2## X is alkyl, aryl, or -NR.sup.5 R.sup.6 R. sup. 5 and R. sup. 6 are each independently hydrogen, alkyl, or is ##STR3## where the dotted line represents an optional double R. sup. 7, and R. sup. 8 and R. sup. 9 are each, independently, hydrogen, alkyl, hydroxy, alkoxy, carbalkoxy, halo, nitro, amino, cyano,

trifluoromethyl, or a carboxylic acid;

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17. 5,049,390, Sep. 17, 1991, Liposome containing immunotherapy agents for treating IgE mediated allergies; Aristo Wojdani, 424/450 []MAGE AVAILABLE]

US PAT NO: 5,049,390 [IMAGE AVAILABLE] L3: 17 of 39

# ABSTRACT:

An immunotherapy agent for the treatment of allergy composed of allergen encapsulated in or covalently bound to a liposome is disclosed. Use of the agent in immunotherapy results in enhanced Ig6 production and reduced IgE production.

5,837,846, Aug. 6, 1991, Indoly1-3 polyamines and their use antagonists of excitatory amino acid neurotransmitters; Nicholas Saccomano, et al., 514/419; 548/495 [IMAGE AVAILABLE]

US PAT NO: 5,037,846 [IMAGE AVAILABLE] L3: 18 of 39

# ABSTRACT:

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This invention relates to certain polyamines found to be present

^\_ venom ^\_ of the Agelenopsis aperta spider. The polyamines of

invention and the salts thereof antagonize excitatory amino acid neurotransmitters, which neurotransmitters affect cells of

organisms, and are useful in antagonizing said neurotransmitters,

in the treatment of excitatory amino acid neurotransmitter mediated

diseases and conditions and in the control of invertebrate pests.

invention also relates to compositions comprising said polyamines salts thereof.

19. 5,034,317, Jul. 23, 1991, Enzyme controlled release system organic conjugate reactant; Michael J. Arnost, et al., 435/18, 4, 6, 14, 19, 21 [IMAGE AVAILABLE]

US PAT NO: 5,034,317 [IMAGE AVAILABLE] L3: 19 of

#### ABSTRACT:

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The present invention provides an enzyme controlled release system and prepared reactant by which an identifiable molecule may be released on demand through the action of an active enzyme. The controlled release system is useful for detection of an analyte of interest present in a test sample in picogram per liter quantities and may be employed in a variety of different modes of use including immunoassays, enzyme

# ligands.

20. 5,821,576, Jun. 4, 1991, 2-Anilino phenylacetic acid derivatives; Amedeo A. Failli, et al., 546/174, 175 [IMAGE AVAILABLE]

amplification systems and the release of pharmacologically active

US PAT NO: 5,021,576 [IMAGE AVAILABLE] L3: 20 of 39

#### ABSTRACT:

There are disclosed compounds of the formula ##STR1## wherein R is hydroxy, lower alkoxy or lower alkoxyamino; R.sup.1 is hydrogen or A(CH.sub.2).sub.n 0--; R.sup.2 is hydrogen or A(CH.sub.2).sub.n 0--, with the proviso that one

of R.sup.1 and R.sup.2 is A(CH.sub.2).sub.n O-- and the other is

hydrogen; n is 1-2;

A is phenoxyethyl, phenoxyphenyl or a group having the formula ##STR2##

X is --N-- or ##STR3## Z is ##STR4## R.sup.3 is hydrogen, lower alkyl

or phenyl; R.sup.4 is hydrogen or lower alkyl; or R.sup.3 and R.sup.4 taken together form a benzene ring; R.sup.5 is hydrogen or lower alkyl; R.sup.6 is hydrogen, halo or lower alkyl;

and the pharmacologically acceptable salts thereof, and their use in the

treatment of inflammatory conditions, such as rheumatoid arthritis,

ulcerative colitis, psoriasis and other immediate hypersensitivity

reactions; in the treatment of leukotriene-mediated naso-bronchial

obstructive air-passageway conditions, such as allergic rhinitis,

المتعادة به محمور مناط cytoprotective agents.

21. 5,013,630, May 7, 1991, Compounds for the cleavage at a position of RNA, oligomers employed for the formation of said compounds, and starting materials for the synthesis of said oligomers; Eiko Ohtsuka, et al., 536/27, 28, 29 [IMAGE AVAILABLE]

L3: 21 of US PAT NO: 5,013,830 [IMAGE AVAILABLE] 39

preferential

ABSTRACT: There is disclosed a compound having a double chain which is an RNA (+chain) and a complementary DNA (-chain), wherein a portion of the DNA (-chain) has been replaced by an RNA or a derivative thereof, and wherein, when the compound is subjected to the action of an enzyme having a ribonuclease Hactivity, it is possible to preferentially RNA (+chain) in a position corresponding to the unsubstituted portion of the DNA (-chain). The compound can thus be used for the

cleavage of a phosphodiester bond in a specific position of RNA. Accordingly, the invention provides a useful means for preparing,

instance, a deletion mutant. There is also disclosed a mixed oligomer which comprises an oligomer of RNA or a derivative thereof and a oligomer, wherein the RNA oligomer or a derivative thereof is conjugated to the DNA oligomer via a phosphate diester linkage between the 51-hydroxyl group and the 31-hydroxyl group in the ribose or deoxyribose moiety. There is further disclosed a nucleoside derivative of a given general formula for use as a starting material.

4,966,712, Oct. 2, 1996, System and method for complement 32. pathway analysis; Argyrios N. Theofilopoulos, et al., 436/501; 435/965, 436/567, 512, 536, 539, 548, 884, 821 [IMAGE AVAILABLE]

L3: 22 of

# ABSTRACT:

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2.1

The present invention relates to systems and methods used to assay for particular complement component fragments. The invention can be used to determine the amount of a particular complement component fragment in a sample. The fragment can be fluid phase or bound to an immune complex. Generally, specific binding agents, such as antibodies, directed to the complement component fragments and immune complement component fragments and immune complement are used in the assay.

23. 4,946,788, Aug. 7, 1998, Purified immunoglobulin-related factor, novel monoclonal antibodies, hybridoma cell lines, processes and applications; Guy Delespesse, 530/388.22; 424/85.8; 435/7.5, 76.21, 172.2, 188, 240.27, 948, 954; 530/388.73, 391.3, 862, 865, 868; 935/95, 100, 102, 103, 104, 106, 107, 108, 110 [IMAGE AVAILABLE]

US PAT NO: 4,946,788 [IMAGE AVAILABLE] L3: 23 of 39

# ABSTRACT:

The invention relates to novel purified human immunoglobulin E binding factors (IgE-BFs), its individual optionally glycosylated proteins, and

monoclonal antibodies to lymphocyte cellular receptors for IgE (Fc. sub..epsilon. R) crossreacting with IgE-Fs, derivatives thereof, processes for the preparation of these antibodies and their derivatives, hybridoma cell lines that produce these antibodies, processes for the preparation of said hybridoma cell lines, the use of the monoclonal antibodies and their derivatives for the qualitative and quantitative determination of IgE-BFs, test kits containing the monoclonal antibodies and/or their derivatives, the use of the monoclonal antibodies for the purification of IgE-BFs, the use of purified IgE-BFs, its individual optionally glycosylated proteins and/or fragments thereof for the

fragments thereof, processes for the purification of IgE-BFs,

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preparations containing them. IgE-BFs and monoclonal antibodies reacting with IgE-BF are important for the diagnosis and therapy of allergic diseases.

24. 4,886,817, Now. 14, 1989, O-functionalized derivatives of substituted isoquinolin-3-ols having cardiotonic and/or phosphodiesterase fraction III inhibiting properties and/or renal vasodilating properties;
Ramesh M. Kanojia, et al., 514/389, 216, 235.2, 253; 544/128, 363; 546/141

US PAT NO: 4,880,817 L3: 24 of 39

### ABSTRACT:

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O-functional derivatives of substituted isoquinolin-3-ol compounds of the general formula ##STRI## that exhibit cardiotonic and/or phosphodiesterase fraction III inhibiting properties and/or renal

vasodilating are pharmacologically active in the treatment of cardiac conditions. Methods for synthesizing and using the compounds are described.

25. 4,867,973, Sep. 19, 1989, Antibody-therapeutic agent conjugates; John H. F. Goers, et al., 424/85.91, 85.8, 86, 87; 514/2, 6, 8;

536/388.7, 388.9, 391.9, 828, 864, 866; 936/10, 22

US PAT NC: 4,867,973 L3: 25 of 39

# ABSTRACT:

This invention relates to antibody-therapeutic agent conjugates having a

therapeutic agent covalently attached to an antibody or antibody fragment. Also described are methods for intermediates in the preparation

of antibody conjugates. Therapeutic in vivo methods utilizing

antibody-therapeutic agent conjugates are described.

26. 4,774,318, Sep. 27, 1988, Snake \_\_venom \_\_ growth arresting peptide; Hans Marquardt, et al., 536/324, 325, 326, 856; 930/10, 260, DIG.821

US PAT NO: 4,774,318

L3: 26 of

#### ARSTRACT:

Novel cytotoxic agents are provided as small polypeptides related to a

low molecular weight peptide derived from Crotalus atrox. The compounds

may be used by themselves or in combination with other reagents, such as

antibodies, for inhibiting cell growth.

27. 4,769,320, Sep. 6, 1988, Immunoassay means and methods useful in

human native prothrombin and human abnormal prothorombin determinations:

Bruce E. Furie, et al., 435/7.92, 7.23, 7.4, 13, 810; 436/69, 536, 548,

868, 811, 815, 825; 538/381, 384, 388.25, 868

US PAT NO: 4,769,320

L3: 27 of

39

# ABSTRACT:

Antibodies which form issume complexes with human native prothrombin

only, in the presence of mixtures of human native prothrombin and human

abnormal prothrombin as well as antibodies which form antibody antigen

complexes with human abnormal prothrombin in the presence of such

mixtures have been obtained. Immunoassay techniques are used for qualitative and quantitative determinations of these antigens in

# human

plasma or serum. Unique methods of obtaining the antibodies are described

including obtaining antibodies to native prothrombin by dissociation of

antigen antibody complexes formed in the presence of calcium ions with a

material having a greater affinity constant for binding with calcium ions

than does prothrowbin. Dissociation of the complex in this manner yields

human native prothrombin antibodies which are specific and non-reactive

with human abnormal prothrombin. A process is described in which assays

are applied to the sensitive detection of vitamin K deficiency and

various forms of liver disease including hepatocellular carcinoma, and to

monitoring of anticoagulant therapy with sodium warfarin. The invention

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Department of Mouth, and Mann, Cr. vines.

20. 4,721,420, Man. 15, 1000, Shabir A\_ samen A\_ garanth arresting puphlics;

Wants Minglands, at al., E20/204; 404/05.01; E14/0, 12; 21; 530/350, 400,

40E, 0E6; 020/10, 20, 200, DIG.821

US PAT NO: 1,721,429 39

13: 20 cf

# RESTERCT:

Novel sytotoxic agents are provided as small polypeptides related

low colocular weight poptide derived from Crotalus atrox. The 001201120

may be used by themselves or in combination with other reagents, בם לשטנ

antibodies, for inhibiting cell growth.

4,710,725, Pag. 20, 1927, Amnitrogen substituted isequinclinel

occapaunds having cardistania, phosphodiesterase fraction III inhibiting

properties and/or renal vasodilating properties; Racosh M. Kanojia, at

al., 514/309; 545/141

UE PAT MB: 4,714,72E 39

13: 29 of

# APETRACT:

Substituted 3-isoquinolinol respected of the general formula

that exhibit cardiotonic and/or phosphodiosteraco fraction III intititing

properties and/or rotal resudilating are pherocoologically active in the

threatment of cerdian conditions. Methods for synthesizing and daing those

coopeunds are described.

20. 4,675,287, Jun. 22, 1987, Monoplomal antibody directed to

genglioside CD. sub.2; Relph N. Roisfeld; st bl., 425/7.23; 42/./25.8;

25,01, 435/7,02, 70,21, 172.2, 240,27, 240, 275, 421/510, 520,

E10, 012; E20,0007 E, 200.0, 200.0E, 201.7, 200, 000, 000; 92E/00, 95, 100;

CO PAT NO. 4, 372, 257 13: 30:45

#### ABSTRACT:

2775

A non-human, mammalian monoclonal receptor produced and secreted by a

hybridosa having the ATCC accession number HB 8568 and methods of

preparing and using same, as well as diagnostics utilizing the receptor

are disclosed. The monoclonal receptor reacts, with cells such as human

neuroectodermal tumors having ganglioside 6D.sub.2 antigen expressed on

their cellular membrane surfaces.

31. 4,572,044, Jun. 9, 1987, Murine monoclonal antibody combining site to human C3b receptor (CR1); Robert D. Schreiber, 436/501; 435/4, 7.21,

7.24, 7.25, 70.21, 172.2, 240.27, 810, 968, 975; 436/504, 506, 507, 512,

518, 536, 540, 548, 815, 821; 935/104, 110

US PAT NO: 4,672,044

L3: 31 of

# ABSTRACT:

A murine monoclonal antibody combining site produced by a hybridoma formed by fusion of cells from a myeloma cell line and lymphocytes that produce antibodies that react (1) with isolated human C3b receptor and

(2) with C3b receptor-bearing cells from a mammal immunized with human C3b receptor is disclosed.

32. 4,671,958, Jun. 9, 1987, Antibody conjugates for the delivery of compounds to target sites; John D. Rodwell, et al., 424/85.91, 1.1, 85.8, 86, 87; 514/2, 6, 8; 530/389.5, 391.5, 391.9, 828, 864, 866 CIMAGE AVAILABLES

US PAT NO: 4,671,958 [IMAGE AVAILABLE] L3: 32 of 39

# ABSTRACT:

A method is described for the covalent attachment of linker groups to specific sites on antibody molecules directed against any desired target antigen (tumor, bacterial, fungal, viral, parasitic etc.). These

can be attached via amide or ester bonds to compounds for delivery which contain available asino or hydroxy groups (e.g., bioactive \_\_cytotoxic agents, dyes, fluors, radioactive compounds, etc.). In addition the linkers can be incorporated into insoluble matrices for use separation schemes which are based upon antibody-antigen interactions. The linkers may be designed so that they are susceptible to cleavage by any one of the serum complement enzymes. When prepared according to the methods described herein, the resulting modified antibody solecule retains the ability to bind antigen and to fix serum complement. when administered to a patient the antibody conjugate binds to

in vivo. As a result of the subsequent activation of the patient's serum complement, the covalently attached compound will be specifically cleaved

at the target site by the proteolytic enzymes of the complement system.

33. 4,661,347, Apr. 28, 1987, Cytotoxic compositions; Hans J. Muller-Eberhard, et al., 424/85.91; 435/188; 530/388.85, 391.7, 391.9, 395, 402, 403, 404, 408

US PAT NO: 4,661,347

L3: 33 of

# ABSTRACT:

A cytotoxic composition is described comprising a moiety having binding affinity to a surface structure of a cell and coupled to a moiety having activity as a structural subunit of C3/C5 convertase.

34. 4,642,284, Feb. 10, 1987, Method and system for detection of

complement pathway activation; Neil Cooper, et al., 435/7.94, 4, 7.4, 28, 965, 966, 971, 975; 436/512, 518, 520, 528, 529, 530, 536, 538, 540, 543, 547, 808, 821

US PAT NO: 4,642,284

L3: 34 of

39

ABCTRACT.

 $\boldsymbol{\mathsf{A}}$  method and system for detecting and preferably measuring the presence

of an activated complement complex in a sample is discussed. The presence

of such an activated complex is indicative of complement pathway activation and includes a first complement component and a second

complement component. The method uses a first binding agent specific to

the first complement component and a second binding agent specific to the

second complement component which when bound with the complex forms an

aggregate. The second specific binding agent includes a label whose

presence is used to detect and measure the amount of aggregate and

therefore activated complex in a sample. An assay system and aggregate

for use in an assay system are also discussed.

35. 4,517,303, May 14, 1985, Specific binding assays utilizing analyte-cytolysin conjugates; J. William Freytag, et al., 436/501; 435/4,

5, 7.21, 7.22, 7.23, 7.25, 7.31, 7.32, 7.5, 7.8, 7.9, 21, 36, 966, 972;

436/512, 526, 541, 803, 813, 815, 817, 827, 828, 829; 930/10, 240, 280,

DIG. 821 [IMAGE AVAILABLE]

US PAT NO: 4,517,303 [IMAGE AVAILABLE]

L3: 35 of

神機 養養 一切

### ABSTRACT:

A novel analyte-cytolysin conjugate and its use in a lipid vesicle

mediated measurement process is described for a wide variety of analytes

present at very low concentration. The method involves forming a reaction

system consisting of analyte, analyte specific binding agent, analyte-cytolysic conjugate, and vesicles containing detectable marker

material in such proportions that uncombined conjugate alters the

permeability of the vesicles resulting in the release and quantitative

detection of marker material which can be correlated with the amount of analyte initially present.

36. 4,454,226, Jun. 12, 1984, Enzymatic immunoassay; Majid Ali, et al.,

531

US PAT NO: 4,454,226

L3: 36 of

\_ABSTRACT:

An enzyme immunoassay for detecting an antigen in a biologic tissue which comprises contacting the fluid or tissue with an antibody specific for the antigen under binding conditions, at least one of the fluid or tissue and antibody having a solid component, contacting resulting solid with a conjugate bindable with the antibody under binding conditions and determining the enzyme activity of the resulting solid phase is described. The conjugate is of peroxidase and an allergen, non-issunoglobulin protein or prisary asino group containing drug an average of 2-3 molecules of peroxidase per molecule of substance with an average molecular weight of about 36,000 daltons, prepared by reacting peroxidase previously treated with phenyl isothiocyanate and form aldehyde groups with the substance to form a Schiff's base which is

titrated with a reducing agent to form a stable conjugate.

37. 4,447,526, May 8, 1984, Homogeneous specific binding assay with carrier matrix incorporating specific binding partner; Patricia A.

Rupchock, et al., 435/7.7; 422/56; 435/7.72, 7.92, 885, 971; 436/528, 539, 535, 537, 816

US PAT NO: 4,447,526

L3: 37 of

39

# ABSTRACT:

producing a

A method for determining the presence of a ligand in, or the ligand binding capacity of a liquid test sample which includes the steps of (a) adding to the sample a conjugate of the ligand and a label, (b) contacting the sample with a test device containing reagents which in conjunction with the conjugate and ligand, are capable of

and the company of the contract of the contract of the contract of

1,566,215, 457 1, 1005, El-brighteshibbe-V-carbiveside-Citis Landy Maria Di, Margara, 20 El., Editor, 20

12: 20 of 12 PRT ND: 4,446,315 30

The present invertion relates to the preparation of and the antituder.

edonosina S'-(kuikydhogen diphrophake) polipound, Elefaderu Elester with

4-carbonacids-2-.tita.-2-ritofuranceyithiacole.

29. 4,430,495, Feb. 7, 1984, Process for preparing lineopypin 8ಗವೆ clindacycin mibanucleotides; Teo E. Matt, et al., E35/16.2, 16.2, 15.4, 16.5, 20, 20

US DOT NO: 4,430,495 13: 39 of

Mayel and useful witch.cleabides of analogs of the well known linecoyain and clindacypin. These mibanuplectides are unexpectedly highly active against Ctroptococcus hecolyticus and Staphylococcus vive. These miborariestides one prepared by using resting cell or

call-from subrands of Stroptonyana rocket, NAME 2527, or extracts of Streptocyces coelicator, MRRL 2522.

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75. 4,897,466, Jan. 30, 1990, Human lymphoblastoid cell line and

hybridowas derived therefrom; James W. Larrick, et al., 424/85.8;

530/809; 935/107

=) t15 100 cit "TL5" IS NOT A RECOGNIZED COMMAND

=) t 15 100 cit

160. 4,772,550, Sep. 20, 1988, Heterogeneous specific binding assay employing an aggregatable binding reagent; Alfred C. Greenquist, 435/7.92, 7.5, 7.93, 7.94, 288, 805, 810; 436/503, 524, 529, 533, 534, 539, 868, 810, 824

=> t 15 125 cit

125. 4,614,793, Sep. 36, 1986, Hepatitis A--subunit antigen; Joseph V. Hughes, et al., 530/350, 418, 806, 826; 930/223

=) t 15 140 cit

140. 4,429,008, Jan. 31, 1984, Thiol reactive liposomes; Frank J.

Martin, et al., 428/402.2; 424/1.1, 85.8, 88, 450; 436/501, 532, 829

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61 ---- Government Interest Text Data
ICLS --- Issue Classification
IN ---- Inventor Information
IPC ---- International Classification
OREF ---- Other Publication References Cited
PARN ---- Parent Case Text Data
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PPDR --- Number of Drawing Sheets
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PPSP ---- Number of Specification Pages
PRIR --- Foreign Application Priority
PTAN ---- PCT Number
PTFD ---- PCT Filing Date
PTFD ---- PCT Publication Date
PTPN ---- PCT Publication Number
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4,429,068, Jan. 31, 1984, Thiol reactive liposomes; Frank

Martin, et al., 428/462.2; 424/1.1, 85.8, 88, 456; 436/561, 532,

140.

# =) t 15 cit 140-152

146. 4,429,668, Jan. 31, 1984, Thiol reactive liposcues; Frank J.

Martin, et al., 428/462.2; 424/1.1, 85.8, 88, 456; 436/501, 532, 829
[IMAGE AVAILABLE]

- 141. 4,421,735, Dec. 20, 1983, Radiolabeled diagnostic compositions and method for making the same; Edgar Haber, et al., 424/1.1, 9; 536/382, 389.1, 389.2, 389.7, 391.5, 392, 404, 405, 408, 409 [IMAGE AVAILABLE]
- 142. 4,466,376, Aug. 23, 1983, Immunological preparations; Arnold R. Sanderson, 424/88, 85.8, 86, 87, 89, 91, 92; 436/543; 530/388.2, 388.73, 389.2, 389.4, 389.5, 389.6, 391.9, 405, 806, 807, 866
- 143. 4,375,972, Mar. 8, 1983, Heterogeneous chemiluminescent immunoassays utilizing metallo porphyrin tag; Peter S. Forgione, et al., 436/531, 546
- 144. 4,368,149, Jan. 11, 1983, Protein hybrid having cytotoxicity and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9;

424/85.91, 88; 536/389.5, 391.7, 866; 935/167

145. 4,366,241, Dec. 28, 1982, Concentrating zone method in heterogeneous immunoassays; Henry K. Tom, et al., 435/7.91; 422/56; 435/5, 7.9, 7.92, 805, 810, 968, 975; 436/541, 800, 807

- 146. 4,363,758, Dec. 14, 1982, Cytotoxic protein hybrid and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9; 424/85.91, 88; 435/172.2; 536/376, 379, 389.6, 391.7, 866; 935/107, 108
- 147. 4,357,273, Nov. 2, 1982, Antitumor protein hybrid and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9; 435/172.2; 514/12; 527/203; 530/389.3, 389.7, 806, 825, 866; 935/107, 108
- 148. 4,350,626, Sep. 21, 1982, Antitumor hybrid and process for the

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402/470.2, 520/209.6, 001, 025/107

140. A, 200, 027, Det. 20, 1901, Pure intravences heren old animal gener general

150. 4,072,526, Jun. 8, 1001, Pullification of Resignable by disciffide innotification; Mooks Echmanaberg, 425/518; 850/300; 425/11, 00; 425/4, 010, 004, 000; 425/513, 546, 000, 000; 520/300.1, 200.0, 201.1, 201.2, 442, 013, 011 [IMAGE GUATLABIE]

151. 4,222,402, Mer. 11, 1980, Reagants and pothod employing channeling; Edward T. Maggio, et al., 435/5, 7.7, 7.91, 068; 425/527, 205

152. A,026,94E, Jul. 19, 1977, Composition and mathed for dublinoining the size and location of systemdial infancts; Edger Mater, 424/1.1; 128/654; 250/303; 424/9 [IMAGE AVAILABLE]

# =) t 1E cit ab 140-1E2

140. 4,420,000, Jan. 21, 1984, Thiol meantive liposcops; Fmant: J.
Martin, et al., 420/400.2; 424/1.1, 85.8, 88, 480; 436/801, 532, 629
IMMOSE EVALUABLES

US PAT NO: 6,480,888 SMARES SVALLABLES L5: 148 of 158

# ARCTRACT:

Liposocco and provided which have a plurality of thick reactive and we

extending outward of the liposocal bilayer. The liposocae forc

co.alent bonds with ligands having thick groups, such as  $\ ^{\prime}$  Fab  $^{\prime}$  .

fragounds. Particularly preferred liposcops include calcidide collectes as

the third committee groups. The third reambire dipensions are usefully

employed in agglutimation essays, such as blood byping and

immibilities, and targeting to openific zells.

144. Ayroly785, Bec. CO, 1982, Radiolabeled diagnostic compositions and

538/382, 389.2, 389.7, 391.5, 392, 484, 485, 488, 489 [IMAGE AVAILABLE]

US PAT NO: 4,421,735 [IMAGE AVAILABLE] L5: 141 of 152

# ABSTRACT:

Protein molecules are covalently bonded to a chelating agent which in turn binds a radioactive molecule. Prior to binding, the radioactive molecule is reduced with dithionite ion and then mixed with protein at a pH of 7.0 to about 8.0.

142. 4,400,376, Aug. 23, 1983, Immunological preparations; Arnold R. Sanderson, 424/88, 85.8, 86, 87, 89, 91, 92; 436/543; 530/388.2, 388.73, 389.2, 389.4, 389.5, 389.6, 391.9, 405, 806, 807, 866

US PAT NO: 4,466,376 L5: 142 of 152

# ABSTRACT:

An immunological preparation of an antigenic material in combination with a major histocompatibility complex antigen, which is itself in the form of complex with a protein with which it is normally

associated in nature or with a modified form of such protein which retains the epitope thereof intact, said antigenic material being attached to the protein of the complex through antibody to that protein, is disclosed as being useful for the production of an immunogenic response in human or veterinary use.

143. 4,375,972, Mar. 8, 1983, Heterogeneous chemiluminescent immunoassays utilizing metallo porphyrin tag; Peter S. Forgione, et al., 436/531, 546

US PAT NO: 4,375,972 L5: 143 of 152

# ABSTRACT:

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A conjugate for use in the detection and quantification of antibodies and

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issuncassay procedures utilizing the conjugate. The conjugate is of reacting with an antigen or an antibody or both and includes a capable of catalyzing a chemiluminescent reaction. The conjugate an antibody or an antigen to which a metallo porphyrin tag is and preferably comprises immunoglobulin to which hemoglobin is attached.

144. 4,368,149, Jan. 11, 1983, Protein hybrid having cytotoxicity and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9; 424/85.91, 98; 530/389.5, 391.7, 856; 935/107

US PAT NO: 4, 368, 149 L5: 144 of

152

# ABSTRACT:

A protein hybrid having cytotoxicity obtained by covalently immunoglobulin or its fragment, which is capable of binding selectively to an antigen possessed by a cell to be destroyed, to a protein,

obtained from Momordica charantia and has an activity to

protein synthesis. This protein hybrid displays remarkable cytotoxicity

against targel cells.

145. 4,366,241, Dec. 28, 1982, Concentrating zone method in heterogeneous immunoassays; Henry K. Tom, et al., 435/7.91; 435/5, 7.9, 7.92, 805, 810, 968, 975; 435/541, 900, 807

4,366,241 L5: 145 of US PAT NO: 152

# ABSTRACT:

Method and apparatus are provided for performing immunoassays

device comprising a relatively small test zone referred to as an immunoscrbing zone, and a relatively large liquid absorbing zone

liquid receiving relationship with said immunosorbing zone. The immunosorbing zone includes a member of an immunological pair ("mip")--ligand and antiligand--bound to a support. A signal producing system is employed in conjunction with said

device

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producing system provides for production of a detectible signal in the immunosorbing zone in relation to the amount of analyte in a sample.

146. 4,363,758, Dec. 14, 1982, Cytotoxic protein hybrid and process for the preparation thereof; Yasuhiko Masuho, et al., 536/391.9; 424/85.91,

88; 435/172.2; 536/376, 379, 389.6, 391.7, 866; 935/107, 188

US PAT NO: 4,363,758

against target cells.

L5: 146 of

ABSTRACT:

A cytotoxic protein hybrid obtained by covalently bonding an immunoglobulin or its fragment, which is capable of linking selectively with an antigen possessed by a cell to be destroyed, to a protein, which is obtained from Phytolacca americana and has an activity to terminate protein synthesis. This protein hybrid displays remarkable cytotoxicity

147. 4,357,273, Nov. 2, 1962, Antitumor protein hybrid and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9; 435/172.2; 514/12; 527/203; 530/389.3, 389.7, 806, 825, 866; 935/107, 108

US PAT NO: 4,357,273

L5: 147 of

05020

ABSTRACT:
Antitumor protein hybrid, composed of a moiety which is substantially the fragment \_^\_ Fab ^\_ of an anti-Lalpha.-fetoprotein antibody and a moiety which is substantially the fragment A of a diphtheria toxin, which is expressed by the following formula (I):

^\_ Fab ^\_ --(S. sub.1 --(X). sub.n --3. sub.2 --FA). sub.s (I)

(where ^\_ Fab ^\_ indicates a moiety which is substantially the fragment \_\_ Fab ^\_ of an anti-.alpha.-fatoprotein antibody; FA indicates a moiety which is substantially the fragment A of diphtheria toxin; X indicates a

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atoms, S.sub.1 indicating a sulfur atom arising from the disulfide bond (--S--S--bond) in an anti-.alpha.-fetoprotein antibody and S.sub.2 a sulfur atom arising from the disulfide bond in a diphtheria toxin; n stands for 6 or 1 and m stands for an integer of 1 to 5). This antitumor protein hybrid has remarkable and specific citotoxicity against tumor cells.

148. 4,350,626, Sep. 21, 1982, Antitumor hybrid and process for the preparation thereof; Yasuhiko Masuho, et al., 530/391.9; 435/172.2; 530/389.6, 866; 935/107

US PAT NO: 4,350,626 L5: 148 of 152

# ABSTRACT:

Antitumor protein hybrid, composed of a moiety which is substantially the fragment ^\_ Fab ^\_ of an antitumor immunogloblin and a moiety which is the subunit A of ricin, which is expressed by the following formula (I):

^\_ Fab ^\_ (S.sub.1 -(X).sub.n -S.sub.2 -RA).sub.m
(I)
(Where \_ ^\_ Fab ^\_ indicates a moiety which is substantially the

# fragment

^\_Fab ^\_ of an antitumor immunogloblin; RA indicates a moiety which is the subunit A of ricin; X indicates a divalent organic radical; S. sub. 1 and S. sub. 2 are both sulfur atoms, S. sub. 1 indicating a sulfur atom arising from the disulfide bond (--S--S-- bond) in an immunoglobulin and S. sub. 2 a sulfur atom arising from the disulfide bond in ricin; n stands for 0 or 1 and m stands for an integer of 1 to 5). This antitumor protein hybrid has remarkable and specific cytotoxicity against tumor cells.

149. 4,296,627, Oct. 26, 1981, Pure intravenous human and animal gamma globulins; Richard M. Condie, 536/396.5; 424/85.5; 536/838

greedins; richard in condite, 330/330.3, 427/63.3, 330/63

US PAT NO: 4,296,027

L5: 149 of

ABSTRACT: Isolated and purified natural, unaltered, undenatured issume

globulin (IgG) for intravenous administration prepared from animal blood

plasma, especially human. The products are characterized by high vield

and high purity. They are unfragmented and unaggregated, i.e., natural

preparation.

150. 4,272,506, Jun. 9, 1981, Purification of reagents by disulfide

immobilization; Moshe Schwarzberg, 435/512; 250/302; 424/1.1, 88; 435/4.

962, 964, 968; 436/513, 546, 866, 825; 536/389.1, 389.3, 391.1, 391.3,

464, 413, 813, 861 [[MAGE AVAILABLE]

US PAT NO: 4,272,586 [IMAGE AVAILABLE] L5: 150 of 152

### ABSTRACT:

A method is provided for preparing immunoassay reagents involving labeled

members of specific binding pairs substantially enriched relative to

contaminating labeled materials. The method involves conjugating a member

of a specific binding pair to a support by a covalent bond which is

cleavable under mild conditions to provide a binding pair

### member-support

conjugate. Combining the binding pair sember-support conjugate with a

labeled composition containing the reciprocal member of the binding pair;

so that the labeled reciprocal, member becomes bound to the support

through the binding of the specific binding pair. Separating the support

to which is bound the labeled member from the remaining labeled material

and then cleaving the bond joining the labeled specific binding pair to

the support to provide labeled reagent for immunoassays. In Agricular,

ntibody is linked to a support by disulfide linkage and a comparison contain the reciprocal antigen to the antibody is labeled with

chromophore, Postalandy Clarecton. The apport is freed of

link cleaved to provide Papeled reagent for immunoassays.

4,233,403, Nov. 11, 1980, Reagents and method employing channeling;

Edward T. Maggie; et al., 435/5, 7.7, 7.91, 968; 435/537, 865

4, 233, 402 US PAT NO:

L5: 151 of

152

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# ABSTRACT:

Method and compositions are provided for chemical analysis of an

which is a member of a specific binding pair of organic substances

consisting of ligand and ligand receptor (antiligand). The method

involves bringing together the following reagents with the analyte in an

aqueous assay medium under mild conditions.

The first reagent is a conjugate of a member of the specific

with a chemical entity which provides a means for chemically

concentration of a compound which acts as a signal mediator. The

reagent is the signal mediator precursor. The third reagent is a conjugate of a member of the specific binding pair with a component of a

signal producing system of which system the signal mediator is a

The amount of signal which can be detected is affected by the

local concentration of the signal mediator. By bringing the reagents in the presence of analyte, where the signal mediator changing means is brought together in a microenvironment with the

conjugated signal producing system component, concentrations of

the signal mediator can be created which differ from the gross concentration of the signal mediator in the assay medium. The

which the signal mediator concentration changing means is in

proximity to the signal producing means in a microenvironment will affect

the observed signal. By appropriate choice of the two conjugates

conjunction with the analyte, the observed signal can be related

amount of analyte in the pains.

specific proportions to substantially optimize the assay sensitivity. The combinations are provided as kits, where ancillary reagents can included, so as to simplify the combination of reagents, as well provide for more accurate measurements and relative proportions reagents.

152. 4,836,945, Jul. 19, 1977, Composition and method for determining the size and location of myocardial infarcts; Edgar Haber, 424/1.1; 128/654; 250/303; 424/9 [IMAGE AVAILABLE]

4,036,945 [IMAGE AVAILABLE] US PAT NO: L5: 152 of 152

# ABSTRACT:

externally

Radioactive labelled antibody for cardiac myosin is injected intravenously after cardiac occlusion and is specifically infarcted myocardium, as are radioactive labelled lower molecular fragments of the antibody such as ( ^\_ Fab ^\_ 1).sub.2, ( ^\_ Fab ^\_ 1), and (Fv). The location and size of the myocardial infarct can be determined measuring the intensity and location of radioactive emission

of the heart.